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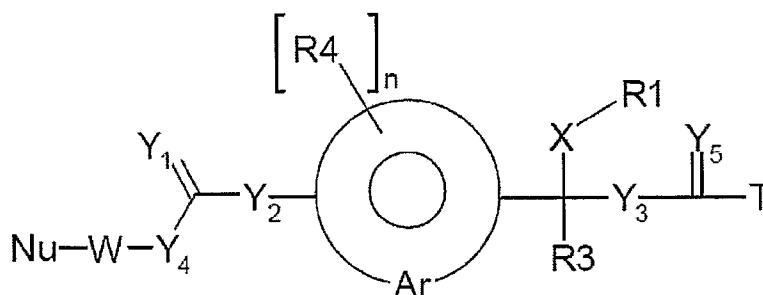
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(54) Title: PRODRUG LINKER



(57) Abstract: A cascade carrier linked prodrug is described which comprises a biologically active moiety and - a masking group having at least one nucleophile and being distinct from the carrier. In one preferred embodiment of the invention, the prodrug has the following structure.

## Prodrug Linker

### Field

- 5 The present invention is directed to polymeric prodrugs having temporary linkages to amino groups of biologically active entities such as peptides, proteins, natural products or synthetic chemical compounds.

### Background

10

Typically, polymers are either used in a non-covalent fashion, with the drug compound physicochemically formulated into a solvent-polymer mixture, or by permanent covalent attachment of a polymer reagent to one of the drug's functional groups.

15

Non-covalent drug encapsulation has been applied to depot formulations for long-acting release profiles. Typically, drug is mixed with polymer material and processed in such fashion, that the drug becomes distributed throughout the bulk polymer material. Such polymer-protein aggregates may be shaped as microparticles which are administered as an injectable suspension or they are formulated as gels which are administered in a single bolus injection. Drug release occurs when the polymer swells or degradation of the polymer allows for diffusion of the drug to the exterior. Such degradation processes may be autohydrolytic or enzyme-catalyzed. An example for a marketed drug based on bolus administration of a drug-polymer gel is Lupron Depot.

20

- 25 An example for a marketed drug based on suspended microparticles is Nutropin Depot.

30

A disadvantage of the non-covalent approach is that in order to prevent uncontrolled, burst-type release of the drug, encapsulation has to be highly efficient by creating a sterically highly crowded environment. Restraining the diffusion of an unbound, water soluble drug molecule requires strong van der Waals contacts, frequently mediated through hydrophobic moieties. Many conformationally sensitive therapeutics such as proteins or peptides are rendered dysfunctional during the encapsulation process and/or during subsequent storage. In addition, such amino-

containing drug compounds readily undergo side reactions with polymer degradation products (D.H. Lee et al., J. Contr. Rel., 2003, 92, 291-299). Furthermore, dependence of the release mechanism upon biodegradation may cause interpatient variability.

- 5 Alternatively, drugs may be conjugated to polymers through permanent covalent bonds. This approach is applied to various classes of molecules, from so-called small molecules, through natural products up to larger proteins.

- Many small molecule medicinal agents, like alkaloids and anti-tumor agents, show  
10 low solubility in aqueous fluids. One way to solubilize these small molecule compounds is to conjugate them to hydrophilic polymers. A variety of water-soluble polymers, such as human serum albumin, dextran, lectins, poly(ethylene glycol) (PEG), poly(styrene-co-maleic anhydride), poly(N-hydroxypropylmethacrylamide), poly(divinyl ether-co-maleic anhydride), hyaluronic acid have been described for this  
15 purpose (R. Duncan, Nature Rev. Drug Disc., 2003, 2, 347-360).

- A major challenge in cancer therapy is to selectively target cytotoxic agents to tumor cells. A promising method to accumulate small molecule anticancer agents in tumor tissue and decrease undesirable side effects of these agents is the attachment of the  
20 cytotoxin to a macromolecular carrier. The passive targeting of polymeric drug conjugates to tumors is based on the so-called enhanced permeability and retention effect (EPR) as described by Matsumura, Y. and Maeda, H., in Cancer Res., 1986, vol 6, pp 6387-6392. As a result, several polymer-drug conjugates have entered clinical trial as anticancer agents.

- 25 Covalent modification of biological molecules with poly(ethylene glycol) has been extensively studied since the late 1970s. So-called PEGylated proteins have shown improved therapeutic efficacy by increasing solubility, reducing immunogenicity, and increasing circulation half-life in vivo due to reduced renal clearance and proteolysis  
30 by enzymes (see, for example, Caliceti P., Veronese F.M., Adv. Drug Deliv. Rev. 2003, 55, 1261-1277).

However, many medicinal agents such as INFalfa2, saquinavir or somatostatin are inactive or show decreased biological activity when a polymer is covalently

conjugated to the drug molecule (T. Peleg-Shulman et al., J. Med. Chem., 2004, 47, 4897-4904).

In order to avoid shortcomings imposed by either non-covalent polymer mixtures or permanent covalent attachment, it may be preferable to employ a prodrug approach for chemical conjugation of drug to polymer carrier. In such polymeric prodrugs, the biologically active moieties are typically linked to the polymeric carrier moiety by a temporary bond formed between the carrier moiety and a hydroxy, amino or carboxy group of the drug molecule (such as is shown in Fig. 1).

Prodrugs are therapeutic agents that are almost inactive per se but are predictably transformed into active metabolites (see B. Testa, J.M. Mayer in *Hydrolysis in Drug and Prodrug Metabolism*, Wiley-VCH, 2003, page 4). The carrier prodrug approach may be applied in such a fashion that the medicinal agent is released in vivo from the polymer in order to regain its biological activity. The reduced biological activity of the prodrug as compared to the released drug is of advantage if a slow or controlled release of the drug is desired. In this case, a relatively large amount of prodrug may be administered without concomitant side effects and the risk of overdosing. Release of the drug occurs over time, thereby reducing the necessity of repeated and frequent administration of the drug.

Prodrug activation may occur by enzymatic or non-enzymatic cleavage of the temporary bond between the carrier and the drug molecule, or a sequential combination of both, i.e. an enzymatic step followed by a non-enzymatic rearrangement. In an enzyme-free in vitro environment such as an aqueous buffer solution, a temporary bond such as an ester or amide may undergo hydrolysis, but the corresponding rate of hydrolysis may be much too slow and not therapeutically useful. In an in vivo environment, esterases or amidases are typically present and may cause significant catalytic acceleration of the kinetics of hydrolysis from twofold up to several orders of magnitude (see, for example, R.B. Greenwald et al. J. Med. Chem. 1999, 42 (18), 3857-3867).

## Definitions based on IUPAC

(as given under <http://www.chem.qmul.ac.uk/iupac/medchem/> (accessed on 8 March 2004))

### 5 Prodrug

A prodrug is any compound that undergoes biotransformation before exhibiting its pharmacological effects. Prodrugs can thus be viewed as drugs containing specialized non-toxic protective groups used in a transient manner to alter or to eliminate undesirable properties in the parent molecule.

10

### Carrier-linked prodrug (Carrier prodrug)

A carrier-linked prodrug is a prodrug that contains a temporary linkage of a given active substance with a transient carrier group that produces improved physicochemical or pharmacokinetic properties and that can be easily removed in vivo, usually by a hydrolytic cleavage. This is shown graphically in Fig. 1.

15

### Cascade prodrug

A cascade prodrug is a carrier prodrug for which the cleavage of the carrier group becomes effective only after unmasking an activating group.

20

### Polymeric cascade prodrug

A polymeric cascade prodrug is a carrier prodrug that contains a temporary linkage of a given active substance with a transient polymeric carrier group for which the cleavage of the carrier becomes effective only after unmasking an activating group.

25

### Bioprecursor prodrug

A bioprecursor prodrug is a prodrug that does not imply the linkage to a carrier group, but results from a molecular modification of the active principle itself. This modification generates a new compound, able to be transformed metabolically or chemically, the resulting compound being the active principle.

30

### Biotransformation

Biotransformation is the chemical conversion of substances by living organisms or enzyme preparations.

Prodrugs fall in two classes, bioprecursors and carrier-linked prodrugs. Bioprecursors do not contain a carrier group and are activated by the metabolic creation of a functional group. In carrier-linked prodrugs the active substance is linked to a carrier moiety by a temporary linkage. This invention is concerned with polymeric carrier-linked or macromolecular prodrugs, where the carrier itself is a macromolecule such as a carrier protein or polysaccharide or polyethylene glycol. Specifically, the invention relates to polymeric carrier-linked prodrugs for which this cleavage between polymer and drug proceeds in two steps according to a cascade mechanism.

Cleavage of a carrier prodrug generates a molecular entity (drug) of increased bioactivity and at least one side product, the carrier. This side product may be biologically inert (for instance PEG) or may have targeting properties (for instance antibodies). After cleavage, the bioactive entity will reveal at least one previously conjugated and thereby protected functional group, and the presence of this group typically contributes to the drug's bioactivity.

In order to implement a prodrug strategy, at least one certain functional group in the drug molecule is employed for attachment of the carrier polymer. Preferred functional groups are hydroxyl or amino groups. Consequently, both the attachment chemistry and hydrolysis conditions vary greatly between these two functionalities.

In a simple one-step mechanism, the prodrug's temporary linkage is characterized by an intrinsic lability or enzyme dependence. The susceptibility of this linkage to hydrolysis in an aqueous environment with or without enzyme catalysis controls the cleavage kinetics between polymeric carrier and drug. Numerous macromolecular prodrugs are described in the literature where the temporary linkage is a labile ester bond. In these cases, the functional group provided by the bioactive entity is either a hydroxyl group or a carboxylic acid (e.g. Y. Luo, MR Ziebell, GD Prestwich, "A Hyaluronic Acid – Taxol Antitumor Bioconjugate Targeted to Cancer Cells", *Biomacromolecules* 2000, 1, 208-218, J Cheng et al, Synthesis of Linear, beta-Cyclodextrin Based Polymers and Their Camptothecin Conjugates, *Bioconjugate Chem.* 2003, 14, 1007-1017, R. Bhatt et al, Synthesis and in Vivo Antitumor Activity of Poly(L-glutamic acid) Conjugates of 20(S)-Camptothecin, *J. Med. Chem.* 2003,

46, 190-193 ; R.B. Greenwald, A. Pendri, C.D. Conover, H. Zhao, Y.H. Choe, A. Martinez, K. Shum, S. Guan, J. Med. Chem., 1999, 42, 3657-3667; B. Testa, J.M: Mayer in Hydrolysis in Drug and Prodrug Metabolism, Wiley-VCH, 2003, Chapter 8).

- 5 Especially for therapeutic biomacromolecules but also for certain small molecule drugs, it may be desirable to link the macromolecular carrier to amino groups of the bioactive entity (i.e. N-terminus or lysine amino groups of proteins). This will be the case if masking the drug's bioactivity requires conjugation of a certain amino group of the bioactive entity, for instance an amino group located in an active center or a
- 10 region or epitope involved in receptor binding. Also, during preparation of the prodrug, amino groups may be more chemoselectively addressed and serve as a better handle for conjugating carrier and drug because of their greater nucleophilicity as compared to hydroxylic or phenolic groups. This is particularly true for proteins which may contain a great variety of different reactive functionalities, where non-
- 15 selective conjugation reactions lead to undesired product mixtures which require extensive characterization or purification and may decrease reaction yield and therapeutic efficiency of the product.

- Amide bonds as well as aliphatic carbamates are much more stable towards hydrolysis
- 20 than ester bonds, and the rate of cleavage would be too slow for therapeutic utility in a carrier-linked prodrug. Therefore it is advantageous to add structural chemical components such as neighbouring groups in order to exert control over the cleavability of the prodrug amide bond. Such additional cleavage-controlling chemical structures that are not provided by the carrier entity nor by the drug are
- 25 called linker. Prodrug linkers can have a strong effect on the rate of hydrolysis of a given temporary bond. Variation of the chemical nature of these linkers allows to engineer the linker properties to a great extent.

- For instance, prodrug linkers may be designed for enzyme-selectivity. Prerequisite for
- 30 enzymatic dependence is that the linker structure displays a structural motif that is recognized as a substrate by a corresponding endogenous enzyme (Fig. 2).

Enzyme-catalyzed acceleration of prodrug cleavage is a desirable feature for organ or cellular targeting applications. Targeted release of the bioactive entity is effected, if

an enzyme, that selectively cleaves the linkage, is specifically present in the organ or cell-type chosen for treatment.

A typical property of an enzyme-dependent temporary linkage is its stability with respect to hydrolysis. The temporary linkage itself will not undergo autohydrolysis at a rate that would release drug to such an extent that a therapeutic effect could be induced in a normal dosing regime. It is only in the presence of the enzyme, that the attack of the enzyme on the linkage causes a significant acceleration of cleavage and concomitant an enhancement of free drug concentration.

10

Several examples have been published for the prodrug activation of amine-containing biologically active moieties by specific enzymes for targeted release. In these cases, cleavage occurs in a one-step process which is catalyzed by the enzyme. G. Cavallaro et al., *Bioconjugate Chem.* 2001, 12, 143-151 describe the enzymatic release of an antitumoral agent by the protease plasmin. Cytarabin is coupled via the tripeptide sequence D-Val-Leu-Lys to the polymer alpha, beta-poly(N-hydroxyethyl)-DL-aspartamide (PHEA). Enzymatic release of cytarabin is effected by the protease plasmin which concentration is relatively high in various kinds of tumor mass.

15

Further examples for antitumoral polymeric prodrugs activated by specific enzymes like beta lactamase (R. Satchi-Fainaro et al., *Bioconjugate Chem.* 2003, 14, 797-804) and cysteine proteases like cathepsin B (R. Duncan et al. *J. Contr. Release* 2001, 74, 135-146) have been described. Wiwattanapatapee et al. (2003) outline a dendrimer prodrug for colonic delivery of 5-aminosalicylic acid. The drug molecule is conjugated by an azo bond to "generation 3" PAMAM dendrimer. 5-aminosalicylic acid is released in the colon by a bacterial enzyme called azo reductase (W. R. Wiwattanapatapee, L. Lomlim, K. Saramunee, *J. Controlled Release*, 2003, 88: 1-9).

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A.J. Garman et al. (A.J. Garman, S.B. Kalindjan, *FEBS Lett.* 1987, 223 (2), 361-365 1987) use PEG5000-maleic anhydride for the reversible modification of amino groups in tissue-type plasminogen activator and urokinase. Regeneration of functional enzyme from PEG-uPA conjugate upon incubation at pH 7.4 buffer by cleavage of the maleamic acid linkage follows first order kinetics with a half-life of 6.1 h. The prodrug cleavage was not investigated in the presence of enzymes, and it can be

30



expected – as explained above - that proteases present in the in vivo environment will significantly contribute to the cleavage of the temporary amide linkage. A further disadvantage of this linkage is the lack of stability of the conjugate at lower pH values. This limits the applicability of the linker to active agents which are stable at  
5 basic pH values, as purification of the active agent polymer conjugate has to be performed under basic conditions to prevent premature prodrug cleavage.

Cascade mechanisms have proven particularly useful in the controlled release of drugs containing amino-group functionalities because linker cleavage characteristics can be  
10 optimized with greater flexibility than in simple one-step prodrugs.

Cascade cleavage is enabled by linker compounds that are composed of a structural combination of a masking group and an activating group. The masking group is attached to the activating group by means of a first temporary linkage such as an ester  
15 or a carbamate. The activating group is attached to an amino-group of the drug molecule through a second temporary linkage, for instance a carbamate. The stability, or susceptibility to hydrolysis of the second temporary linkage is dependent on the presence or absence of the masking group. In the presence of the masking group, the second temporary linkage is highly stable and unlikely to release drug with  
20 therapeutically useful kinetics. In the absence of the masking group, this linkage becomes highly labile, causing rapid cleavage and drug release.

Cleavage of the first temporary linkage is the rate-limiting step in the cascade mechanism. This first step may induce a molecular rearrangement of the activating  
25 group such as a 1,6-elimination. The rearrangement renders the second temporary linkage so much more labile that its cleavage is induced. Ideally, the cleavage rate of the first temporary linkage is identical to the desired release rate for the drug molecule in a given therapeutic scenario. Furthermore, it is desirable that cleavage of the second temporary linkage is instantaneous after its lability has been induced by cleavage of  
30 the first temporary bond.

A variety of examples exist for cascade carrier prodrugs where the masking group functionality is performed by the carrier polymer itself as shown diagrammatically in Fig. 3. In the systems discussed below, the masking group is not only part of the

carrier but has also been engineered for enzyme-dependence (Fig. 4). Only in the presence of a corresponding enzyme is the rate of cleavage of the first temporary linkage sufficiently accelerated for therapeutic use.

- 5 R.B. Greenwald, A. Pendri, C.D. Conover, H. Zhao, Y.H. Choe, A. Martinez, K. Shum, S. Guan, J. Med. Chem., 1999, 42, 3657-3667 & PCT Patent Application WO-A-99/30727 describe a methodology for synthesizing poly(ethylene glycol) prodrugs of amino-containing small molecule compounds based on 1,4- or 1,6-benzyl elimination. In this approach, poly(ethylene glycol) as the polymeric carrier is  
10 attached to the benzyl group by means of a first temporary linkage such as an ester, carbonate, carbamate, or amide bond. The benzyl group serves as the activating group, and the PEG polymer also has the function of the masking group in this cascade cleavage mechanism. The amino group of the drug molecule is linked via a second temporary linkage, containing a carbamate group, to the benzyl moiety. The  
15 release of PEG from the drug molecule is initiated by enzymatic cleavage of the first temporary linkage followed by a rapid 1,4- or 1,6-benzyl elimination, initiating cleavage of the second temporary linkage.

- The same linker system is also used for releasable poly(ethylene glycol) conjugates of  
20 proteins (S. Lee, R.B. Greenwald et al. Bioconj. Chem. 2001, 12 (2), 163-169). Lysozyme is used as model protein because it loses its activity when PEGylation takes place on the epsilon-amino group of lysine residues. Various amounts of PEG linker were conjugated to the protein. Regeneration of native protein from the PEG conjugates occurs by enzymatic cleavage in rat plasma or in non-physiological high  
25 pH buffer.

- Greenwald et al. published in 2000 a poly(ethylene glycol) drug delivery system of amino-containing prodrugs based on trimethyl lock lactonization (R.B. Greenwald et al. J.Med.Chem. 2000, 43(3), 457-487; PCT Patent Application No. WO-A-  
30 02/089789). In this prodrug system, substituted o-hydroxyphenyl-dimethylpropionic acid is linked to PEG by an ester, carbonate, or carbamate group as a first temporary linkage and to amino groups of drug molecules by means of an amide bond as second temporary linkage. The rate-determining step in drug release is the enzymatic

cleavage of the first linkage. This step is followed by fast amide cleavage by lactonization, liberating a potentially toxic aromatic lactone side product.

Similar prodrug systems were described by F.M.H. DeGroot et al. (WO02083180 and  
5 WO04043493A1) and D. Shabat et al. (WO04019993A1). WO02083180 discloses a prodrug system with elongated and multiple linkers based on 1,(4+2n) elimination. The masking moieties in these examples were specifically designed for enzymatic cleavage. This approach was extended to dendritic prodrug system where one enzymatic activating event triggered the release of more than one drug molecule  
10 (WO04043493A1). WO04019993A1 discloses a similar prodrug system based on a self-immolative dendrimer releasing many drug moieties upon a single enzymatic activating event. These systems are characterized by the absence of a polymeric carrier. Instead, oligomerization of prodrug linker components provides for a high molecular weight of the prodrug, and prodrug cleavage generates linker residues and  
15 free drug, but no polymeric entity is released.

The disadvantage in the abovementioned prodrug systems described by Greenwald, DeGroot and Shabat is the release of potentially toxic aromatic small molecule side products like quinone methides after cleavage of the temporary linkage. The  
20 potentially toxic entities are released in a 1:1 stoichiometry with the drug and can assume high in vivo concentrations. This risk factor is even greater if self-immolative dendritic structures based on oligomers of the activating group are employed and more aromatic side products than drug molecules are released.

25 More recently, R.B. Greenwald et al. (Greenwald et al. J. Med.Chem. 2004, 47, 726-734) described a PEG prodrug system based on bis-(N-2-hydroxyethyl)glycin amide (bicin amide) linker. In this system two PEG molecules are linked to a bicin molecule coupled to an amino group of the drug molecule. The first two steps in prodrug activation is the enzymatic cleavage of both PEG molecules. Different linkages  
30 between PEG and bicin are described resulting in different prodrug activation kinetics. The main disadvantage of this system is the slow hydrolysis rate of bicin amide conjugated to the drug molecule ( $t_{1/2} = 3$  h in phosphate buffer) which results in the release of a bicin-modified prodrug intermediate that may show different

pharmacokinetic and pharmacodynamic properties as compared to the parent drug molecule.

5 Cascade prodrugs with masking groups that are part of the carrier polymer are limited in the control of drug release kinetics. As masking group cleavage is the rate-limiting step in the cascade mechanism, its molecular structure governs the kinetics. If the carrier polymer is identical to the masking group, the structural flexibility is restricted to the polymers' features. Alternatively, if the polymer requires structural modification in order to match the requirements for controlled cleavage, synthesis of  
10 corresponding structures may become more difficult. Also, the incorporation of masking group features into a polymer may change its safety profile.

Therefore is it preferred to structurally separate the masking group and the carrier. This may be achieved by employing a permanent bond between polymer carrier and  
15 activating group. This stable bond does not participate in the cascade cleavage mechanism. If the carrier is not serving as a masking group and the activating group is coupled to the carrier by means of a stable bond, release of potentially toxic side products such as the activating group is avoided. The stable attachment of activating group and polymer also suppresses the release of drug-linker intermediates with  
20 undefined pharmacology.

Systems have been developed for targeted delivery of therapeutic agents by rendering the masking group enzyme-dependent. Only in the presence of a corresponding enzyme is the rate of cleavage of the first temporary linkage connecting the masking  
25 group with the activating group sufficiently accelerated for therapeutic use.

Antczak et al. (Bioorg Med Chem 9 (2001) 2843-48) describe a reagent which forms the basis for a macromolecular cascade prodrug system for amine-containing drug molecules. In this approach an antibody serves as carrier, a stable bond connects the  
30 antibody to an activating moiety, carrying an enzymatically cleavable masking group. Upon enzymatic removal of the ester-linked masking group, a second temporary bond cleaves and releases the drug compound, as shown in Fig. 6.

D. Shabat et al. (Chem. Eur. J. 2004, 10, 2626-2634) describe a polymeric prodrug system based on a mandelic acid activating moiety. In this system the masking group is linked to the activating moiety by a carbamate bond. The activating moiety is conjugated permanently to a polyacrylamide polymer via an amide bond. After  
5 enzymatic activation of the masking group by a catalytic antibody, the masking group is cleaved by cyclization and the drug is released. The activating moiety is still connected to the polyacrylamide polymer after drug release.

M.-R. Lee et al. describe (Angew. Chem. 2004, 116, 1707-1710) a similar prodrug  
10 system based on mandelic acid activating moiety and an enzymatically cleavable ester-linked masking group.

In all of these described prodrug-polymer systems the masking group is specifically designed to be substrate to an enzyme, and masking group cleavage will almost  
15 entirely depend upon enzymatic catalysis with the disadvantages of interpatient variability, injection site variability and poor in vitro-in vivo correlation.

A major drawback of predominantly enzymatic cleavage is interpatient variability. Enzyme levels may differ significantly between individuals resulting in biological  
20 variation of prodrug activation by enzymatic cleavage. Enzyme levels may also vary depending on the site of administration, for instance it is known that in the case of subcutaneous injection, certain areas of the body yield more predictable therapeutic effects than others. To reduce this unpredictable effect, non-enzymatic cleavage or intramolecular catalysis is of particular interest (see, for example, B. Testa, J.M:  
25 Mayer in Hydrolysis in Drug and Prodrug Metabolism, Wiley-VCH, 2003, page 5).

Furthermore, it is difficult to establish an in vivo-in vitro correlation of the pharmacokinetic properties for such enzyme-dependent carrier-linked prodrugs. In the  
absence of a sound in vivo-in vitro correlation the optimization a release profile  
30 becomes a cumbersome task.

Also, the need for enzyme selectivity imposes a severe limitation on the structural features that can be used in the prodrug linker. This restriction greatly hinders the

development of a sound structure-activity relationship and consequently the optimization of linker cleavage kinetics.

- For these reasons, there is a need to provide novel linker and/or carrier technologies
- 5 for forming polymeric prodrugs of amine containing active agents in order to overcome the limitations of the described polymeric prodrugs.

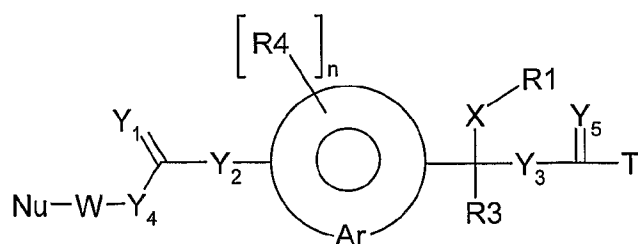
Detailed description of the invention

The present invention addresses the disadvantages described above. The invention provides for polymeric cascade prodrugs characterized by a masking group containing  
5 a nucleophile and being distinct from the carrier.

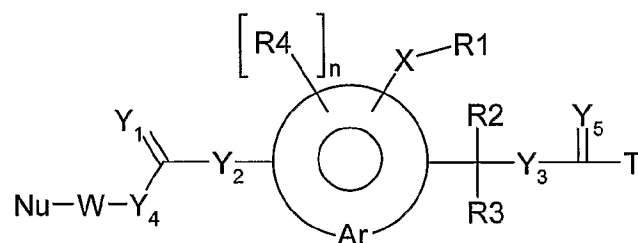
The nucleophile is in a suitable distance to a first temporary linkage with an aromatic activating group capable of undergoing a 1,(4+2p) elimination reaction (with  $p = 0, 1, 2, 3, 4, \dots$ ) after cleavage of the first temporary linkage. The invention is further  
10 characterized by the activating group being connected to the amino group of a drug molecule through a second temporary bond which is cleaved as a consequence of the 1,(4+2p) elimination. An additional structural feature is the attachment of a polymeric carrier to the activating group by means of a permanent bond.

15 The masking groups according to the present invention contain at least one nucleophile Nu. This nucleophile, which can be for example a primary, secondary or tertiary amino group can assist the cleavage of the masking group from the activating moiety by intramolecular catalysis or cyclization.

The invention provides for polymeric cascade prodrugs and corresponding polymeric cascade prodrug linker reagents of Formula Ia or Ib.



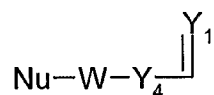
Formula Ia



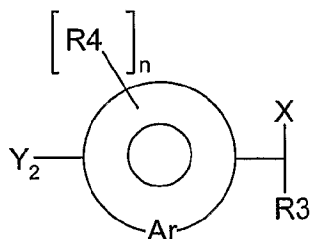
Formula Ib

wherein Y<sub>1</sub> to Y<sub>5</sub>, R<sub>1</sub> to R<sub>4</sub>, T, X, W, Nu and Ar are defined below:

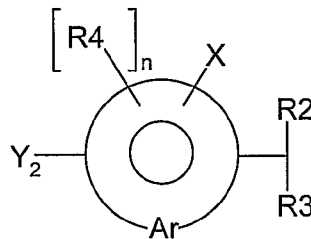
Native drug release is effected by a two step mechanism. The first step is the rate-determining cleavage of the first temporary linkage between the masking group



and the activating moiety



or

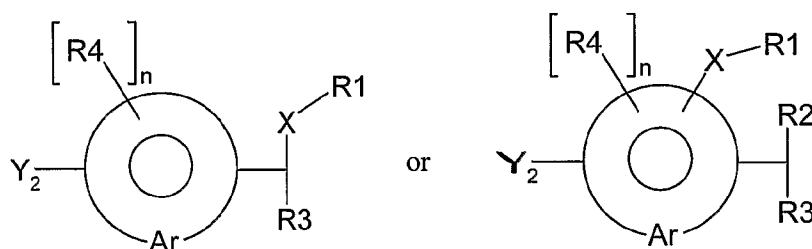




from the polymeric prodrug in vivo.

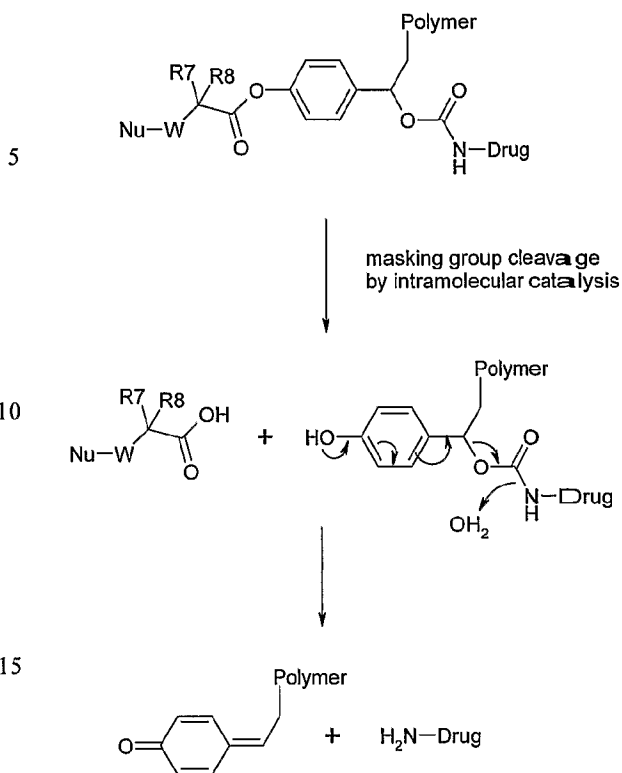
As described above, cleavage of the masking group may be mediated by an enzymatic or a non-enzymatic step, such as pH-dependent hydrolysis or intramolecular cyclization. In the preferred embodiment of the invention, cleavage is effected non-enzymatically by intramolecular cyclization or catalysis. The half-life of the cleavage kinetics in an aqueous buffer of pH 7.4 at 37 °C of the masking group according to the present invention is preferably between 1 hour and 6 months, more preferably between 1 day and 3 months, and most preferably between 1 day and 2 months.

The second and final step in release of the regenerated native drug is the fast, spontaneous and irreversible so-called 1,4- or 1,6 or 1,(4+2p) (in which p = 2, 3, 4 or higher) elimination of the



moiety of the remaining polymeric prodrug of formula Ia or formula Ib, respectively.

This mechanism of native drug release from a polymeric prodrug triggered by hydrolytic cleavage of the masking group followed by a 1,6-elimination step of the activating group is exemplified by a polymeric prodrug according to the present invention.



20 **Definition of Y<sub>1</sub> to Y<sub>5</sub>, R<sub>1</sub> to R<sub>4</sub>, T, X, W, Nu and Ar in formula Ia or Ib**

T is D or A

In the case where the inventive structure is a polymeric cascade prodrug linker reagent, T is A, and A is a leaving group. Non-limiting examples of suitable leaving groups A include but are not limited to chloride, bromide, fluoride, nitrophenoxy, imidazolyl, N-hydroxysuccinimidyl, N-hydroxybenzotriazolyl, N-hydroxyazobenzotriazolyl, pentafluorophenoxy, N-hydroxysulfosuccinimidyl, or any other leaving group known by those skilled in the art.

30 In the case where the inventive structure is a polymeric cascade prodrug, T is D, and D is a residue of an amine-containing biologically active material including but not limited to small molecule bioactive agents or biopolymers like proteins, polypeptides and oligonucleotides (RNA, DNA), peptide nucleic acids (PNA),

Note that in this description reference is often made to prodrugs. A true prodrug is found when T is the residue of the amine-containing biologically active material or moiety. If T is a leaving group A then the formula represents a polymeric cascade prodrug linker reagent. For simplicity these will be referred to prodrugs in this description. It will be understood from the context whether a true prodrug or a reagent as a precursor is meant.

Suitable organic small molecule bioactive moieties include, without limitation, moieties such as central nervous system-active agents, anti-infective, anti-neoplastic, antibacterial, anti-fungal, analgesic, contraceptive, anti-inflammatory, steroidal, vasodilating, vasoconstricting, and cardiovascular agents with at least one primary or secondary amino group. Non-exclusive examples of such compounds are daunorubicin, doxorubicin, idarubicin, mitoxantron, amino glutethimide, amantadine, diphenylsulfon, ethambutol, sulfadiazin, sulfamerazin, sulfamethoxazol, sulfalen, clinafloxacin, moxifloxacin, ciprofloxacin, enoxacin, norfloxacin, neomycin B, spectinomycin, kanamycin A, meropenem, dopamin, dobutamin, lisinopril, serotonin, carbutamid, acivicin, etc.

Suitable proteins and polypeptides having at least one free amino group include but are not limited to ACTH, adenosine deaminase, agalsidase, albumin, alfa-1 antitrypsin (AAT), alfa-1 proteinase inhibitor (API), alteplase, anistreplase, anicod serine protease, antibodies (monoclonal or polyclonal, and fragments or fusions), antithrombin III, antitrypsins, aprotinin, asparaginases, biphalin, bone-morphogenic proteins, calcitonin (salmon), collagenase, DNase, endorphins, enfuvirtide, enkephalins, erythropoietins, factor VIIa, factor VIII, factor VIIa, factor IX, fibrinolysin, fusion proteins, follicle-stimulating hormones, granulocyte colony stimulating factor (G-CSF), galactosidase, glucagon, glucagon-like peptides like GLP-1, glucocerebrosidase, granulocyte macrophage colony stimulating factor (GM-CSF), phospholipase-activating protein (PLAP), gonadotropin chorionic (hCG), hemoglobins, hepatitis B vaccines, hirudin, hyaluronidases, iduronidase, immune globulins, influenza vaccines, interleukins (1 alfa, 1 beta, 2, 3, 4, 6, 10, 11, 12), IL-1 receptor antagonist (rhIL-1ra), insulins, interferons (alfa 2a, alfa 2b, alfa 2c, beta 1a, beta 1b, gamma 1a, gamma 1b), keratinocyte growth factor (KGF), transforming growth factors, lactase, leuprolide, levothyroxine, luteinizing hormone, lyme vaccine,

natriuretic peptide, pancrelipase, papain, parathyroid hormone, PDGF, pepsin, platelet activating factor acetylhydrolase (PAF-AH), prolactin, protein C, octreotide, secretin, sermorelin, superoxide dismutase (SOD), somatropins (growth hormone), somatostatin, streptokinase, sucrase, tetanus toxin fragment, tilactase, thrombins, thymosin, thyroid stimulating hormone, thyrotropin, tumor necrosis factor (TNF), TNF receptor-IgG Fc, tissue plasminogen activator (tPA), TSH, urate oxidase, urokinase, vaccines, plant proteins such as lectins and ricins.

Also included herein is any synthetic polypeptide or any portion of a polypeptide with in vivo bioactivity. Furthermore, proteins prepared by recombinant DNA methodologies including mutant versions of aforementioned proteins, antibody fragments, single chain binding proteins, catalytic antibodies and fusion proteins are included.

Preferred proteins are antibodies, calcitonin, G-CSF, GM-CSF, erythropoietins, hemoglobins, interleukins, insulins, interferons, SOD, somatropin, TNF, TNF-receptor-IgG Fc, and GLP-1.

X is a spacer moiety such as R5-Y6

20

Y<sub>1</sub>, Y<sub>2</sub> can each be either O, S, or NR6, independently of each other.

Y<sub>3</sub>, Y<sub>5</sub> can each be either O or S, independently of each other.

Y<sub>4</sub> is O, NR6, or -C(R7)(R8)-

Y<sub>6</sub> is O, S, NR6, succinimide, maleimide, unsaturated carbon-carbon bonds or any heteroatom containing a free electron pair, or is not present.

25

R2 and R3 are selected independently from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryl, cyano, nitro, halogen, carboxy, carboxyalkyl, alkylcarbonyl, carboxamidoalkyl, etc.

30

The term "heteroalkyl" in the context of the present invention denotes (linear, cyclical or branched) alkyl chains where the alkyl chains contain or are substituted with at any position one or more heteroatoms, selected independently from O, S, N, P, Si, Cl, F, Br, I, etc. or groups, selected independently from carboxamide, carboxylic ester,

phosphonate ester, phosphate ester, double or triple bonds, carbamate, urea, thiourea, thiocarbamate, oxime, cyano, carboxyl, carbonyl, etc.

Each R<sub>4</sub> substitution on Ar may be the same or different and is selected independently  
 5 from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl, substituted or non-substituted linear, branched, or cyclical alkoxy, substituted or non-substituted linear, branched, or cyclical heteroalkyloxy, aryloxy, heteroaryloxy, cyano, halogen, etc.

10

R<sub>4</sub> is selected preferably from small substituents such as hydrogen, methyl, ethyl, ethoxy, methoxy, and other C<sub>1</sub> to C<sub>6</sub> linear, cyclical or branched alkyls and heteroalkyls.

15 n is zero or a positive integer.

R<sub>7</sub> and R<sub>8</sub> are selected independently from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryl, cyano, halogen, etc.

20

R<sub>5</sub> is selected from substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryl, etc.

R<sub>6</sub> is selected from hydrogen, substituted or non-substituted linear, branched or  
 25 cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryl, etc.

R<sub>1</sub> is a polymer.

Non-limiting examples for suitable polymers are polyalkyloxy-based polymers like  
 30 poly(propylene glycol) or poly(ethylene glycol), dextran, chitosan, hyaluronic acid and derivatives, alginate, xylan, mannan, carrageenan, agarose, cellulose, starch, hydroxyethyl starch (HES) and other carbohydrate-based polymers, poly(vinyl alcohols), poly(oxazolines), poly(anhydrides), poly(ortho esters), poly(carbonates), poly(urethanes), poly(acrylic acids), poly(acrylamides) such as

poly(hydroxypropylmethacrylamide) (HMPA), poly(acrylates), poly(methacrylates) like poly(hydroxyethylmethacrylate), poly(organophosphazenes), poly(siloxanes), poly(vinylpyrrolidone), poly(cyanoacrylates), poly(esters) such as poly(lactic acid) or poly(glycolic acids), poly(iminocarbonates), poly(amino acids) such as poly(glutamic acid), collagen, gelatin, copolymers, grafted copolymers, cross-linked polymers, hydrogels, and block copolymers from the above listed polymers.

Hydrogels may be defined as three-dimensional, hydrophilic or amphiphilic polymeric networks imbibing large quantities of water. The networks are composed of homopolymers or copolymers, are insoluble due to the presence of covalent chemical or physical (ionic, hydrophobic interactions, entanglements) crosslinks. The crosslinks provide the network structure and physical integrity. Hydrogels exhibit a thermodynamic compatibility with water which allows them to swell in aqueous media.(see.: N.A. Peppas, P. Bures, W. Leobandung, H. Ichikawa, Hydrogels in pharmaceutical formulations, Eur. J. Pharm. Biopharm. 2000, 50, 27-46). The chains of the network are connected in such a fashion that pores exist and that a substantial fraction of these pores are of dimensions of between 1 and 1000 nm. By selecting certain polymerization conditions, the hydrogel may be obtained in the form of an amorphous gel or as beaded resin. Such soft beads may have a diameter of between 1 and 1000 micrometer.

Hydrogels may be synthesized from the polymers and copolymers listed above and physically cross-linked or chemically cross-linked by radical, anionic or cationic polymerization, by chemical reactions like condensation or addition reactions as described in W.E. Hennink and C.F. van Nostrum, Adv. Drug Del. Rev. 2002, 54, 13-36.

Further examples include branched and hyperbranched polymers. Examples for such polymers include dendrimers and other dense star polymers. (R. Esfand, D.A. Tomalia, Drug Discov Today, 2001, 6(8), 427-436; P.M. Heegaard, U. Boas, Chem. Soc. Rev. 2004 (33(1), 43-63; S.M. Grayson, J.M. Frechet, Chem. Rev. 2001, 101 (12), 3819-3868).

R1 can also be a biopolymer like a protein. Non-limiting examples of such polymers include albumin, antibodies, fibrin, casein, and other plasma proteins.

Each R1 polymer can carry one or more biologically active substances linked to the polymer by conjugation with a second prodrug linker as described herein or any other linker known to the person skilled in the art. The polymers may have further substituents and may be functionalized for attachment to the spacer moiety X. Non-limiting examples of such functional groups comprise carboxylic acid and activated derivatives, amino, maleimide, thiol, sulfonic acid and derivatives, carbonate and derivatives, carbamate and derivatives, hydroxyl, aldehyde, ketone, hydrazine, isocyanate, isothiocyanate, phosphoric acid and derivatives, phosphonic acid and derivatives, haloacetyl, alkyl halides, acryloyl, arylating agents like aryl fluorides, hydroxylamine, disulfides like pyridyl disulfide, vinyl sulfone, vinyl ketone, diazoalkanes, diazoacetyl compounds, epoxide, oxirane, and aziridine.

Preferred functional groups for the R1 polymer include but are not limited to thiol, maleimide, amino, carboxylic acid and derivatives, carbonate and derivatives, carbamate and derivatives, aldehyde, and haloacetyl.

Especially preferred functional groups include thiol, maleimide, amino, carboxylic acid and derivatives, carbamate and derivatives, and carbonate and derivatives thereof.

Non-limiting examples for suitable bonds or groups formed between X and R1 include disulfide, S-succinimido, amide, amino, carboxylic ester, sulfonamide, carbamate, carbonate, ether, oxime, hydrazone, urea, thiourea, phosphate, phosphonate, etc.

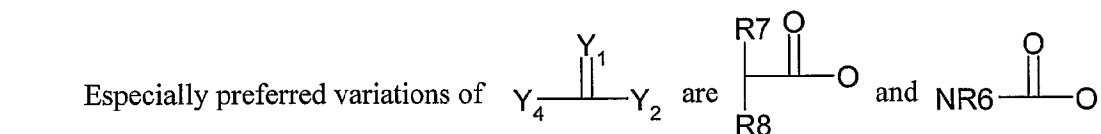
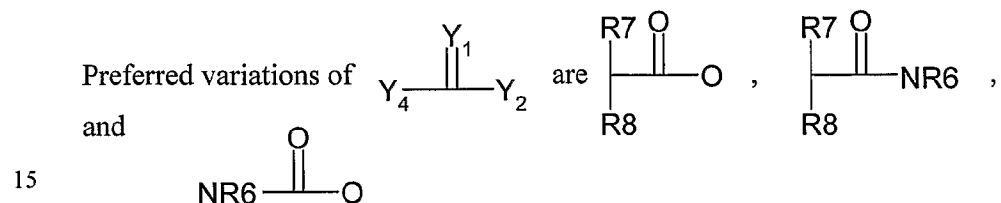
Preferred bonds or groups formed between X and R1 comprise S-succinimido, amide, carbamate, and urea.

Preferably, the R1 polymers are well hydrated, degradable or excretable, nontoxic and non-immunogenic in mammals. Preferred R1 polymers include polyalkoxy-based polymers like polyethylene glycol and polyethylene glycol reagents as those

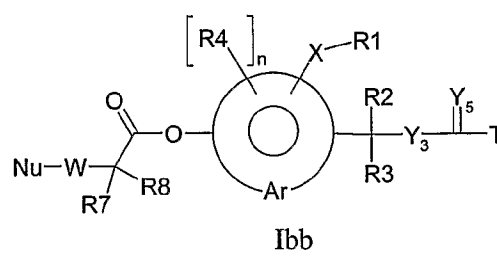
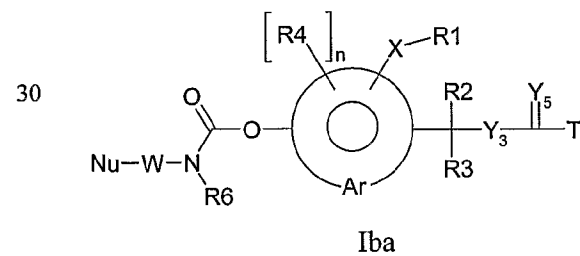
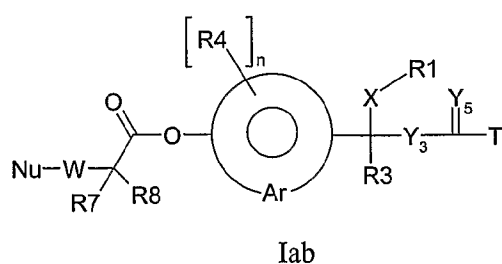
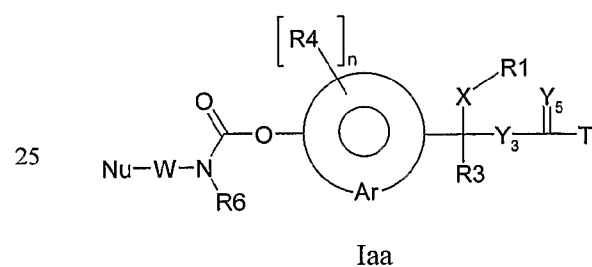
described in Nektar Inc. 2003 catalog "Nektar Molecule Engineering – Polyethylene Glycol and Derivatives for Advanced PEGylation" and branched, hyperbranched, cross-linked polymers and hydrogels, and proteins like albumin.

- 5 W is selected from substituted or non-substituted linear, branched or cyclical alkyl, aryls, substituted aryls, substituted or non-substituted linear, branched or cyclical heteroalkyl, substituted or nonsubstituted heteroaryls, etc.

- W is selected preferably from non-toxic substituted or non-substituted linear,  
10 branched or cyclical alkyls or heteroalkyls.



- 20 forming polymeric prodrugs of the following formulas:





In formula Iaa and Iba R6 may also be Nu-W.

At least one Nu is present in Nu-W.

- 5 Nu is a nucleophile that can perform a nucleophilic attack at the carbonyl carbon of  

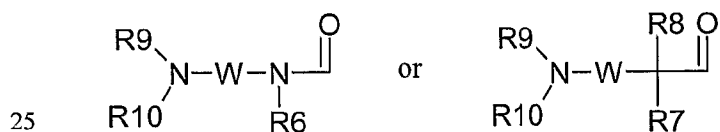
$$\text{Y}_4 - \overset{\text{Y}_1}{\parallel} - \text{Y}_2$$
 and thus catalyse the cleavage of the masking group by intramolecular  
 catalysis or cyclization (figure 8). Figure 8 shows an example according to formula Ia  
 or Ib wherein the cleavage of the masking group is by intramolecular cyclization. In  
 cases where Nu only catalyses the cleavage of the masking group by intramolecular  
 10 catalysis, no cyclical product of the masking group is formed.

Preferred nucleophiles include primary, secondary and tertiary amino groups, thiol,  
 carboxylic acid, hydroxylamine, hydrazine, and nitrogen containing heteroaryl.

- Especially preferred nucleophiles include primary, secondary and tertiary amino  
 groups. In order to effectively catalyse the cleavage of the masking group, the spacing  
 15 between the nucleophile Nu and Y<sub>2</sub> is preferably between three and fifteen atoms.  
 More preferably, the spacing between Nu and Y<sub>2</sub> is between four and ten atoms. The  
 at least one nucleophile Nu may be attached anywhere to W (e.g. at the terminus or in  
 the middle of W) or may be part of W.

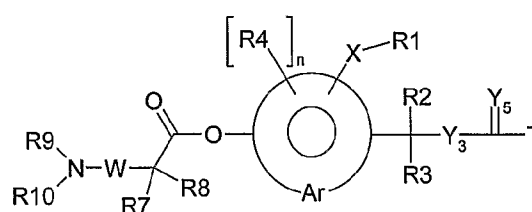
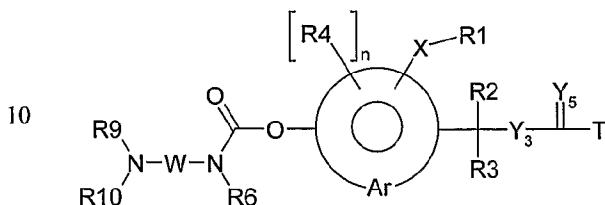
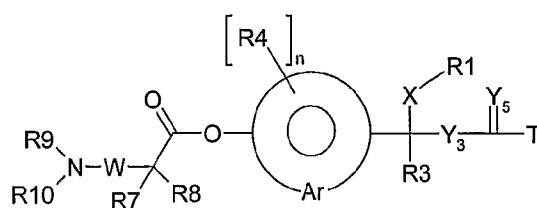
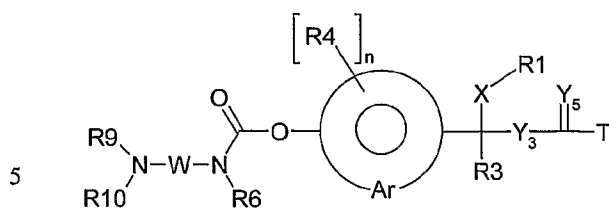
- 20 Preferred variations for the masking group  $\text{Nu}-\text{W}-\overset{\text{Y}_1}{\parallel}_{\text{Y}_4}$

are selected independently from



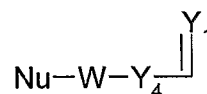
where  $\begin{array}{c} \text{R9} \\ \diagdown \\ \text{N}-\text{W} \\ \diagup \\ \text{R10} \end{array}$  forms a primary, secondary or tertiary amine nucleophile Nu.

- These preferred variations result in polymeric prodrugs according to the following  
 30 formulas:



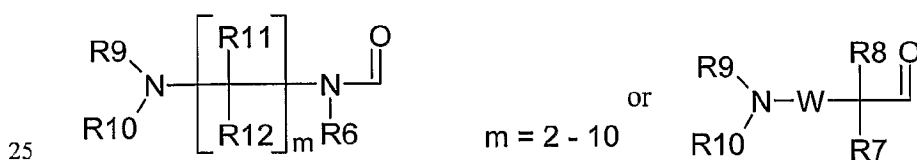
- 15 R9, R10 are selected independently from hydrogen, substituted or non-substituted alkyl or heteroalkyl, substituted or non-substituted aryl or heteroaryl.

Especially preferred variations for the masking group

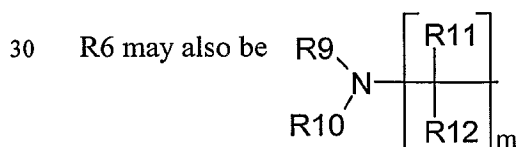


20

are selected from



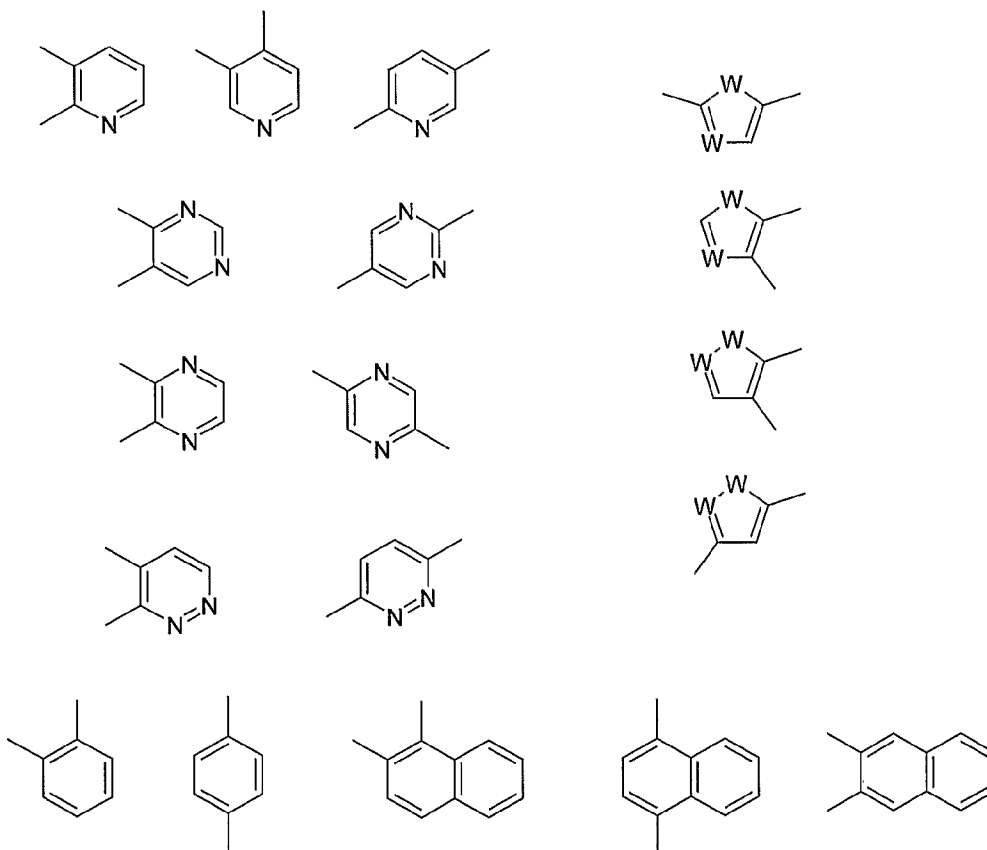
Preferably, R9, R10, R11 and R12 are selected independently from hydrogen, substituted or non-substituted alkyl and R7 and/or R8 are not hydrogen.



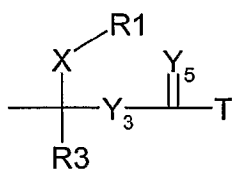
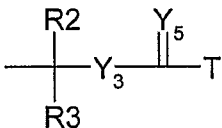
and is preferably not hydrogen.

Surprisingly it was found, that the masking group can modify irreversibly the amine containing biologically active moiety when the nucleophile Nu is absent in the masking group. As shown in the example section, during release of the bioactive moiety insulin from a polymeric prodrug with a pentanoyl masking group which is not part of the present invention (as it does not contain a nucleophile), approximately 30% of the insulin molecule was modified with the masking group by acyl transfer. The mechanism of an example of this modification where D contains an additional free amino group that serve as nucleophile for acyl transfer from the masking group is shown in figure 9.

Ar of formula Ia or Ib is a multi-substituted aromatic hydrocarbon or a multi-substituted aromatic heterocycle. To be aromatic, the number of pi electrons must satisfy the Hückel rule ( $4n+2$ ) and the cycle has to be planar. A huge variety of compounds satisfy these criteria and thus are suitable as Ar in formula Ia or Ib. Non-limiting preferred aromatic moieties include:



wherein W is O, N, or S, independent from each other.

5  $\text{Y}_2$  and  in formula Ia or  $\text{Y}_2$  and  in formula Ib have to be arranged on the aromatic ring in such a fashion that a 1,4- or 1,6- or 1,(4+2p), with p= 2, 3, 4 and higher, elimination can take place (see above). For example, in the case of a 6-membered ring, the substituents have to be arranged *ortho* or *para*.

10

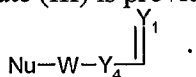
Preferred moieties for Ar are mono- and dicyclic aromatic hydrocarbons or aromatic heterocycles.

Especially preferred moieties are monocyclic five- or six-membered aromatic  
15 hydrocarbons or aromatic heterocycles.

### General synthesis procedures of the polymeric prodrugs

Synthesis of representative examples of polymeric prodrugs according to the present  
20 invention is described in the Examples section.

Prodrugs of the present invention can be prepared in various different fashions. Figure 10 shows general routes for the synthesis of the polymeric prodrugs of the present invention according to formula Ia.

25 In a first method, intermediate (III) is provided by acylating  $\text{Y}_2$  of starting material (II) with the masking group .

For this, X or Nu may have to be protected with a reversible protecting group  $\text{PG}_1$ . Suitable protecting groups are described in TW Greene, P.G.M. Wuts, Protective  
30 groups in organic synthesis, 1999, John Wiley & Sons, 3<sup>rd</sup> ed.. From intermediate (III) two alternative routes can be used to yield (Iaa). In a first route intermediate (III) is activated by an activating agent such as 4-nitrophenyl chloroformate or disuccinyl carbonate to yield (IV). The amine containing drug molecule is attached to (IV) to yield (V) by displacing the leaving group of the activated intermediate (IV). After

deprotection of X such as by treating intermediate (V) with reagents like trifluoroacetic acid or DTT (where applicable) deprotected intermediate (V) is then reacted with polymer R1 to yield the polymeric prodrug (Iaa).

- 5 In a second route the polymer R1 is first attached to the intermediate (III) after deprotection of X (where applicable) to form intermediate (VI). After an activation step intermediate (VII) is formed. (VII) is reacted with the amine containing drug molecule to form the polymeric prodrug (Iac).
- 10 In a second method, intermediate (VIII) is provided by activating starting material (II) by an activating agent such as 4-nitrophenyl chlorofomate. For this, Y<sub>2</sub> and/or X may have to be protected with a protecting group PG<sub>2</sub> and/or PG<sub>1</sub>. Amine containing drug is reacted with intermediate (VIII) to form (IX). In a first route, Y<sub>2</sub> of (IX) is selectively deprotected and acylated to form intermediate (V) which is further
- 15 processed to (Iaa) as described above. In a second route X is selectively deprotected and reacted with polymer R1 to form intermediate (X). Y<sub>2</sub> of (X) is then deprotected and acylated to form the polymeric prodrug (Iac).

In a third method starting material (II) is reacted with polymer R1 to form

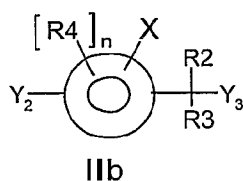
- 20 intermediate (XI). In one route, intermediate (XI) can be acylated to form intermediate (VI) which processed as described above to form polymeric prodrug (Iaa). In a second route, Y<sub>2</sub> is protected by the protecting group PG<sub>2</sub>, activated and reacted with the amine containing drug molecule to form (X). Intermediate (X) is then processed as described above to form the polymeric prodrug (Iac).

25

For all methods described, further functional groups such as Y<sub>3</sub> or nucleophiles present in Nu-W may have to be protected with suitable protecting groups.

Polymeric prodrugs according to formula Ib can be prepared by methods described

- 30 above for prodrugs according to formula Ia using starting material IIb instead of II in figure 10.



It is understood, that linker structures according to the outlined invention and carrying protecting groups or leaving groups as described and used in the synthesis of corresponding polymeric prodrugs are considered within the range of the invention.

5

### **Application of the polymeric prodrugs in molecular therapy**

For polymeric cascade prodrugs it is desirable for the cleavage kinetics of the first temporary linkage to proceed under conditions present in the blood circulation of the human body (pH 7.4, 37°C). Most importantly, cleavage of the first temporary linkage should be based on hydrolysis and exhibit none or only very limited dependence upon chemical or biochemical or physicochemical entities present in the human blood circulation such as enzymes, salts or binding proteins.

15 It was now surprisingly found that the rate of cleavage of the first temporary linkage connecting the masking group with the activating group as well as its dependence upon blood components can be controlled by neighbouring group effects mediated by nucleophilic functional groups (such as primary, secondary or tertiary amines) present in the masking group and positioned in a distance to the corresponding temporary  
20 bond. If the masking group is structured in this fashion, an intramolecular reaction with contribution of the nucleophile governs the kinetics of the linkage (Fig. 5 and Fig. 8).

Key advantage of the polymeric prodrugs of the present invention is their  
25 predominantly non-enzymatic cleavage: the half-life of the prodrug in suitably buffered human blood plasma of pH 7.4 (with aqueous buffer concentration < 50 %) is at least 50 % of the half-life of the prodrug in enzyme-free buffer pH 7.4.

This feature allows for better predictability and control of release rates after  
30 administration to a living organism and reduces interpatient variability.

In contrast to the enzymatic dependency of masking group removal as described in abovementioned examples of Antczak et al., Shabat et al. and Lee et al., a higher

level of control over release rates can be achieved if the masking group has enzyme-independent, self-eliminating properties.

The masking groups according to the present invention contain at least one  
5 nucleophile Nu. Structural features of this masking group such as nucleophilicity of the amine group and ring-forming capacity may be systematically optimized in order to precisely adjust the rate of prodrug cleavage. Such intramolecular reactions resulting in unmasking and subsequent rearrangement are highly independent from enzymes due to the fact that intramolecular reactions are generally preferred over  
10 intermolecular reactions as shown diagrammatically in Fig. 8.

In another embodiment of the invention, independency of prodrug cleavage from enzyme levels is achieved by providing a prodrug containing a sterically demanding carrier group as is shown in Fig. 7.

15 Such encapsulation or sterical protection by the sterically demanding carrier group may be conferred by a branched, hyperbranched, crosslinked or self-assembled structure of the carrier polymer. Such polymers tend to form a densely packed molecular volume, as exemplified for instance in dendrimers, dense star polymers or  
20 bead-shaped nano- and microparticles or amorphous gels. If the linkage of the polymer carrier to the drug is located in the interior of the polymer carrier, the linked drug will be efficiently encapsulated and protected from enzymatic attack. In this case, sterical hindrance by the polymer prevents enzymes from accessing and cleaving the temporary linkages.

25 In yet another embodiment, enzyme-independent prodrug cleavage is achieved by combining an intramolecular self-eliminating masking group with an encapsulating hyperbranched or crosslinked or self-assembled carrier.

30 A further advantage of the present invention is the release of an unmodified biologically active moiety. In cases where the biologically active moiety contains further reactive functional groups like amino groups of lysine residues in proteins, an unwanted side reaction between the masking group and the biologically active moiety can occur. The reactive functional groups of the biologically active moiety may react

with the masking group, forming a stable covalent bond and resulting in the release of a modified biologically active moiety. This potential side reaction is shown schematically in Figure 9. The occurrence of such side reactions is shown in the examples section using polymeric prodrugs which are not part of the present invention with simple masking groups like a pentanoyl residue without a nucleophile Nu present in the masking group and as described by Antczak et al. or Lee et al. The side reaction in this linker system is suppressed using polymeric prodrugs according to the present invention with intramolecularly activated masking groups that contain nucleophiles Nu (see example section).

Enzyme-independent release control enables depot formulations without the need for encapsulation. Until now, many biocompatible materials like hydrogels with large pore sizes could not be used for depot formulations due to their lack of encapsulation properties. From such well-hydrated and mechanically soft biocompatible materials, biologically active moiety would be released too fast for most therapeutic applications. In combination with the prodrug linkers described in this invention, the carrier material may be optimized for its biocompatibility properties as the release is solely governed by the linker cleavage kinetics and does not require chemical or enzymatic degradation of the polymer carrier itself.

Release rates are governed by a substantially non-enzymatic chemical reaction which is in turn dependent on the molecular structure of the linker. Systematic or random modifications of the chemical structure, for instance by changing substituents in one or more positions, for instance a masking group in a cascade prodrug, allows for the generation of prodrug linkers with differing release rates. It is therefore possible to create a variety of prodrug linkers and select those fast or slow cleaving prodrug linkers according to the demands posed by a given medicinal or therapeutic application.

Another advantageous feature which is part of this invention is the attachment of the polymer carrier through a stable covalent bond to an activating moiety involved in a double or cascade prodrug release mechanism. As part of this invention, the activating moiety remains attached to the polymer carrier after drug release and therefore cannot diffuse into the environment. Permanent linkage of the polymer carrier to the activating



group greatly reduces the any side-reactivity of the activating moiety and the probability of unwanted toxic effects. In other polymeric cascade prodrugs known in the art, the activating moiety is released in addition to the drug. Due to the molecular rearrangement mechanisms used in cascade prodrugs, the activating moiety is released  
5 in a highly reactive form and may cause direct damage to surrounding biomolecules, or potentially toxic derivatives of the activating moiety may be formed in vivo.

### Description of the Figures

- 10 Fig. 1 shows a carrier-linked prodrug.  
Fig. 2 shows an enzyme-dependent carrier-linked prodrug.  
Fig. 3 shows a cascade prodrug where the masking group is part of the carrier.  
Fig. 4 shows an enzyme-dependent cascade prodrug where the masking group is part of the carrier.  
15 Fig. 5 shows a self-cleaving cascade prodrug where the masking group is separate from the carrier.  
Fig. 6 shows an enzyme-dependent cascade prodrug where the masking group is separate from the carrier.  
Fig. 7 shows a cascade prodrug where the carrier is sterically protecting the masking  
20 group.  
Fig. 8 shows cleavage of the masking group by intramolecular cyclisation.  
Fig. 9 shows a possible side reaction of polymeric prodrug activation.  
Fig.10 shows general synthesis methods.  
Fig.11 shows mass spectra of prodrug released insulin molecules.

## Examples

### Materials

Fmoc-amino acids, resins and PyBOP were purchased from Novabiochem and are  
5 named according to the catalogue. Fmoc-Ado-OH was obtained from Neosystem. All  
additional chemicals were purchased from Sigma Aldrich. Recombinant human  
insulin was from ICN Biomedicals (USA). Maleimide-PEG5k was obtained from  
Nektar (USA). 5-(and-6)-carboxyfluorescein succinimidyl ester (mixed isomers) was  
obtained from Molecular Probes.

10

### Solid phase synthesis reaction medium

Solid phase synthesis was performed on NovaSyn TG Sieber amide resin with a  
loading of 0.17 mmol/g or 2-chlorotrityl chloride resin with a loading of 1.4 mmol/g.  
Syringes equipped with polypropylene frits were used as reaction vessels.

15

### Standard coupling cycle for fmoc-protected amino acids

For fmoc protecting-group removal, the resin was repeatedly (three times, 4 min each)  
agitated with 2/2/96 (v/v/v) piperidine/DBU/DMF and repeatedly (six times) washed  
with DMF.

20 Coupling of fmoc-protected amino acids to free amino groups on resin was achieved  
by agitating the resin with 3 equivalents (eq) of fmoc-amino acid, 3 eq PyBOP and 6  
eq DIEA in relation to free amino groups in DMF for 60 min.  
Finally, the resin was repeatedly (five times) washed with DMF.

### Standard cleavage protocol for TentaGel Sieber amide resin

Upon completed synthesis, the resin was washed with DCM, dried in vacuo and  
treated repeatedly (five times) with 97/2/1 (v/v) DCM/TES/TFA. After evaporation,  
compounds were purified by preparative RP-HPLC (Waters 600).

### Standard cleavage protocol for 2-chlorotrityl chloride resin

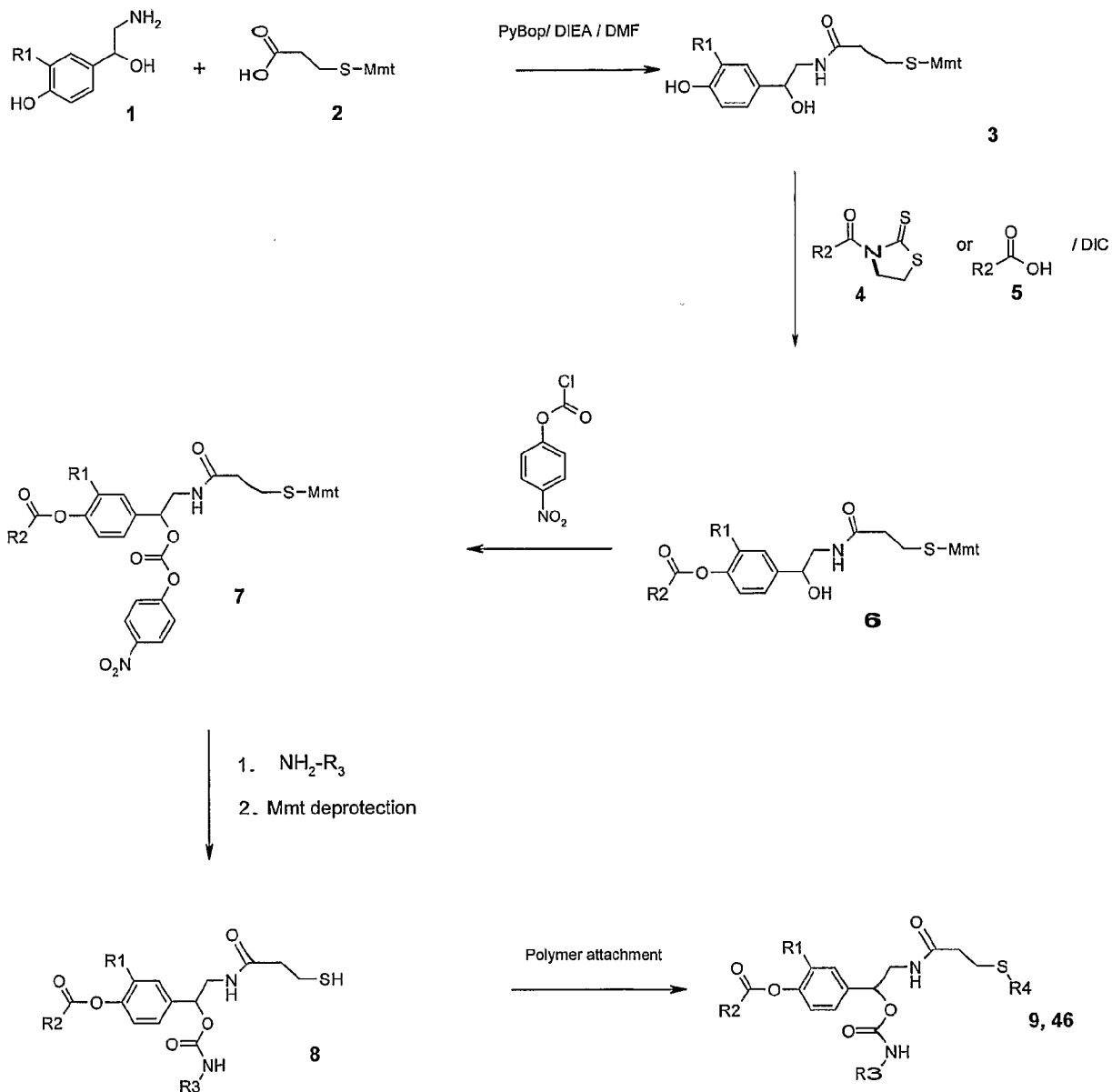
30 Upon completed synthesis, the resin was washed with DCM, dried in vacuo and  
treated two times for 30 minutes with 65/35 (v/v) HFIP/DCM. After combining the  
eluates the volatile components were evaporated.

Analysis

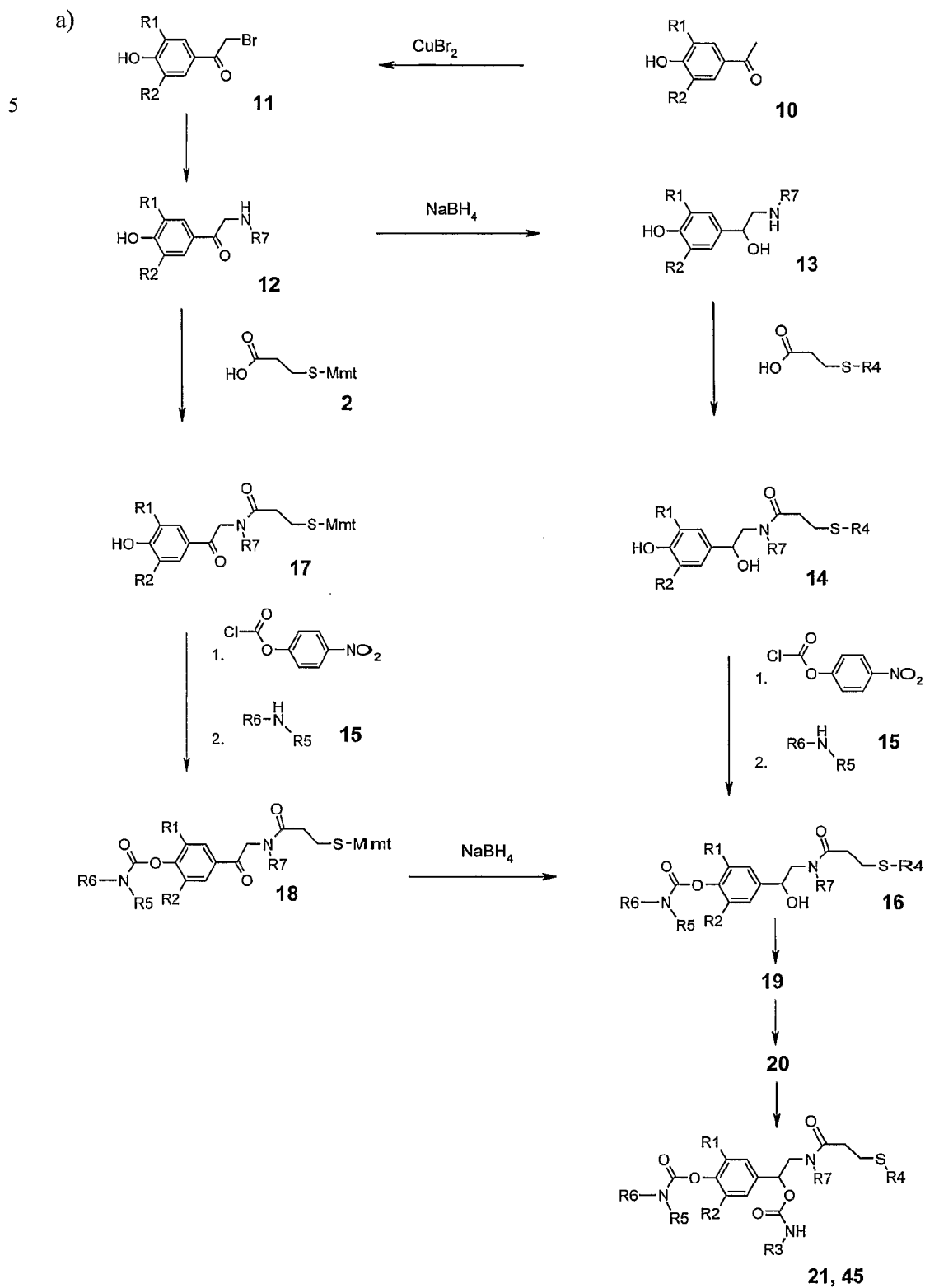
Mass spectrometry (MS) was performed on a Waters ZQ 4000 ESI instrument and spectra were, if necessary, interpreted by Waters software MaxEnt.

- 5 Size exclusion chromatography was performed using an Amersham Bioscience AEKTA basic system equipped with a Superdex 200 column (Amersham Bioscience). NMR spectra were recorded on a Bruker AC300.

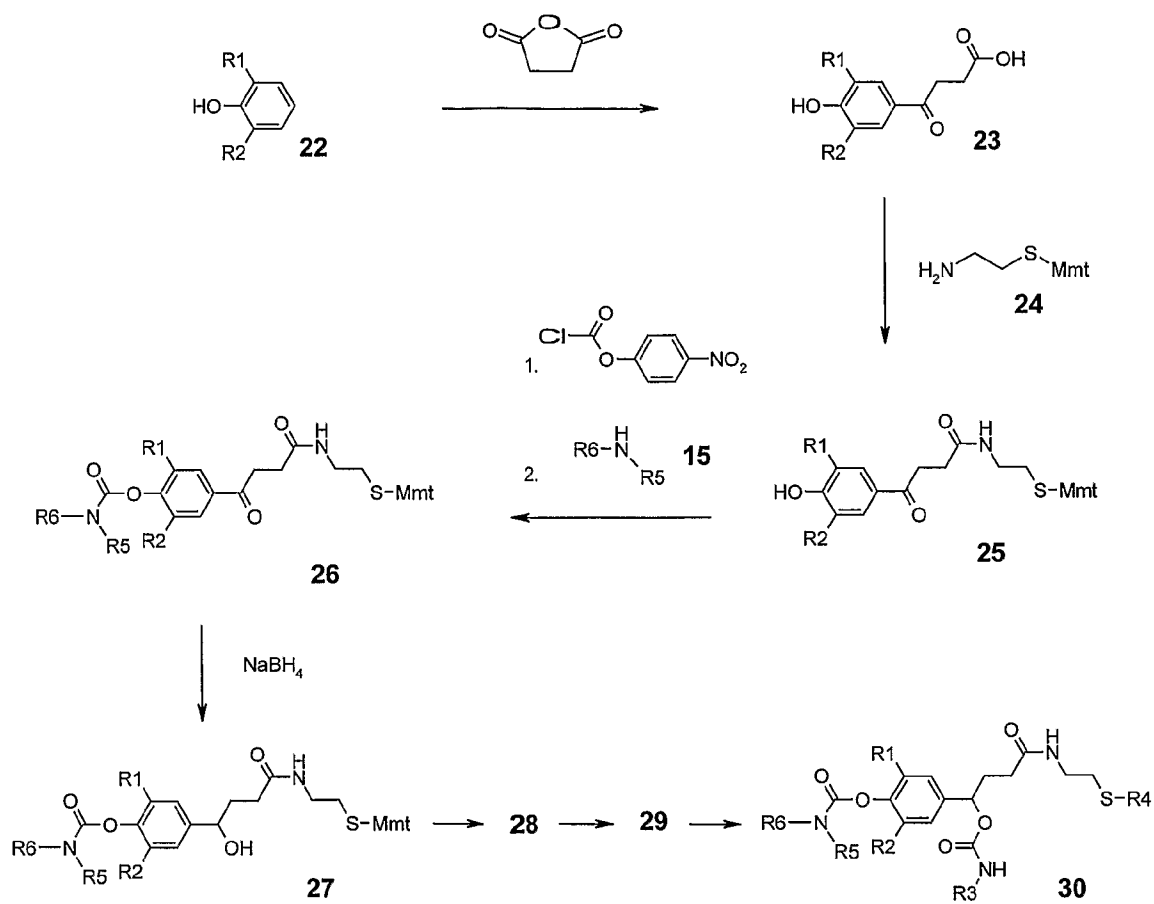
10 Overview - synthesis of polymeric prodrugs according to formula Ia with ester-linked masking groups



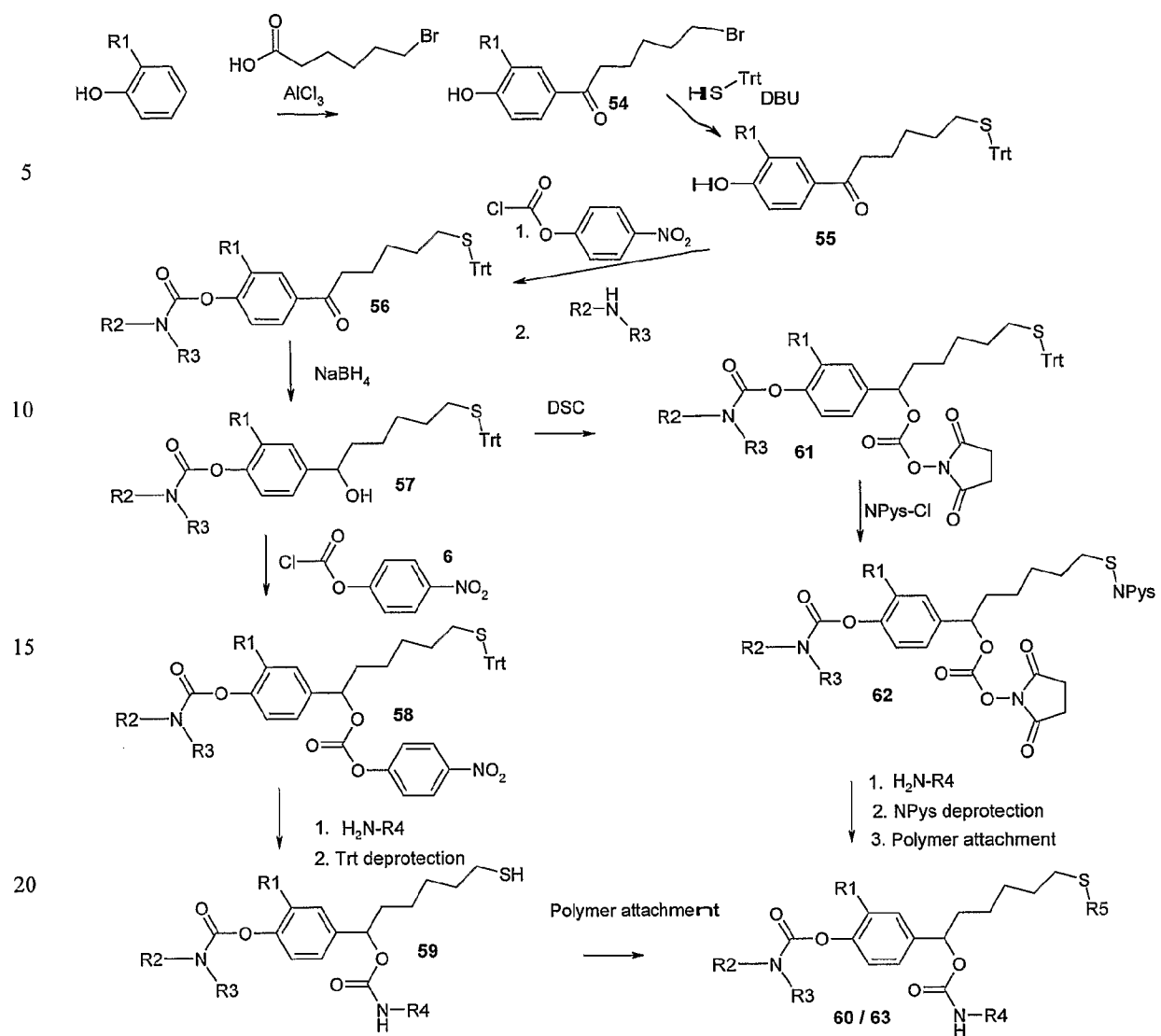
Synthesis of polymeric prodrugs according to formula Ia with carbamate-linked masking groups



b)

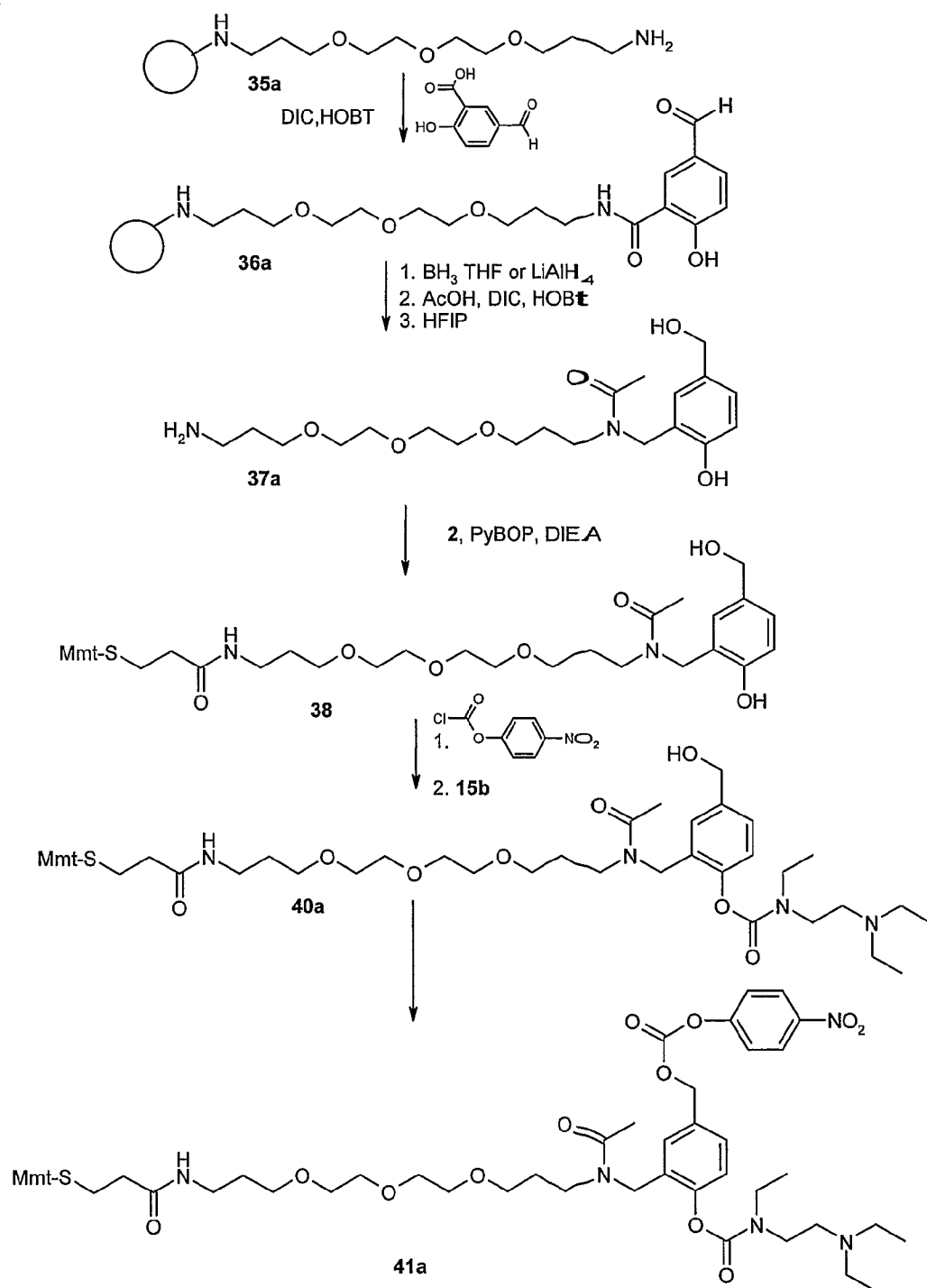


c)

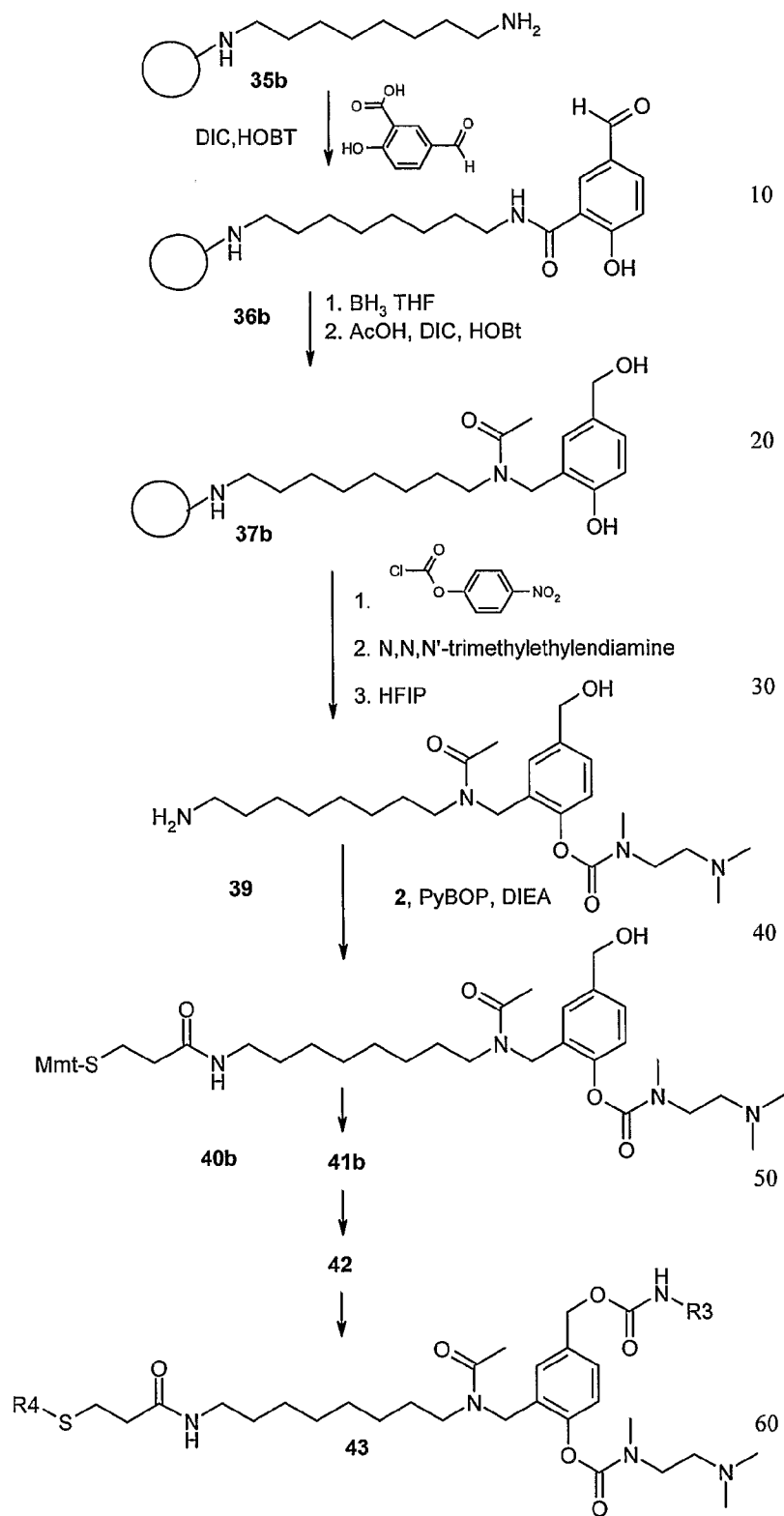


Synthesis of polymeric prodrugs according to formula IIb with a carbamate-linked masking group

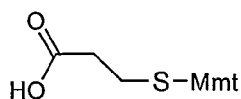
a)



50 b)





Synthesis of compound 2

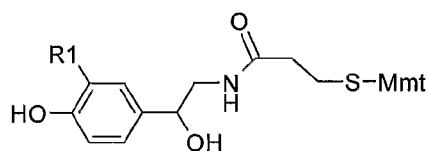
5

**2**

Mmt-chloride (1 eq) and mercaptopropionic acid (1.1 eq) were dissolved in TFA and incubated for 30 min. The solvent was removed under reduced pressure. The product was dissolved in pyridine, diluted in water, acidified by acetic acid and extracted with  
 10 ether. The ether phase was separated and dried over Na<sub>2</sub>SO<sub>4</sub>. Solvent was removed under reduced pressure and product **2** was purified by RP-HPLC.

Synthesis of compounds 3a and 3b

15

**3a** R1 = H20 **3b** R1 = OMe

Octopamine hydrochloride (**1a**) (2 eq), DIEA (4 eq), and PyBOP (1 eq) were dissolved in DMF, **2** (1eq) was added and the mixture was reacted for 50 min at room temperature. After addition of acetic acid (7 eq) product **3a** was purified by RP-HPLC.

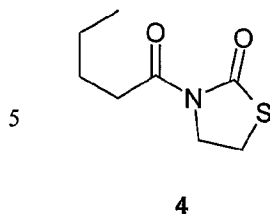
25 **3b** was synthesized from normetanephrine hydrochloride (**1b**) as described above.

**3a**: MS [M+Na]<sup>+</sup> = 536 (MW+Na calculated = 536.2 g/mol)

**3b**: MS [M+Na]<sup>+</sup> = 566 (MW+Na calculated = 566.2 g/mol)

30

### Synthesis of mercaptothiazolide **4**

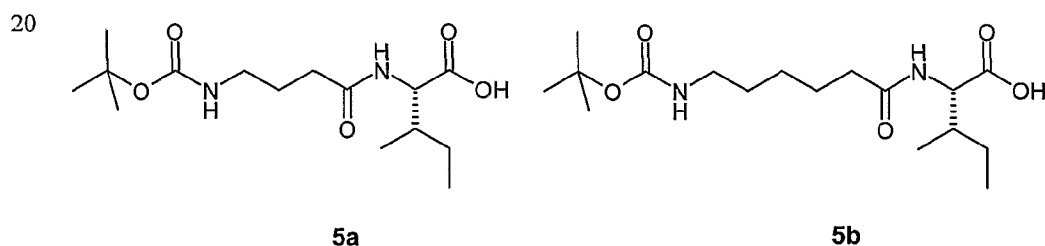


2-Mercaptothiazoline and triethylamine (1.5 eq) were dissolved in dry THF and pentanoyl chloride (1 eq) was added. The mixture was stirred for 1 h at 50 °C under inert gas atmosphere and was allowed to cool to room temperature. 0.5 N aqueous HCl was added and the separated organic phases were dried over Na<sub>2</sub>SO<sub>4</sub>. After concentration *in vacuo* the residue was purified by silica gel column chromatography using heptane/ethyl acetate (1/1) as mobile phase. Mercaptothiazolide **4** was collected as a viscous yellow oil.

15

**4** R<sub>f</sub> (heptane/ethyl acetate 1:1) = 0.7

### Synthesis of compounds **5a** and **5b**



25

General synthesis protocol:

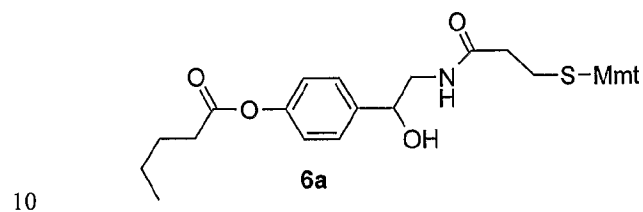
1 g 2-chlorotriethyl chloride resin (loading 1.6 mmol/g) was incubated for 1 h with 850 mg (2.4 mmol) Fmoc-Ile-OH and 840 µl (4.8 mmol) DIEA in 5 ml 1/1 DCM/DMF. After fmoc removal and washing of the resin with DMF, boc-aminobutyric acid was coupled to 0.5 g resin according to the standard coupling method. Compound **5a** was cleaved from the resin with 97/1/2 (v/v) DCM/TFA/TES for 45 min. After neutralisation with pyridine, solvents were removed under reduced pressure and **5a** was purified by RP-HPLC.

**5b** was synthesized from boc-aminohexanoic acid as described above.

**5a** MS  $[M+Na]^+ = 339.2$  (MW+Na calculated = 339.4 g/mol)

**5b** MS  $[M+Na]^+ = 367.4$  (MW+Na calculated = 367.5 g/mol)

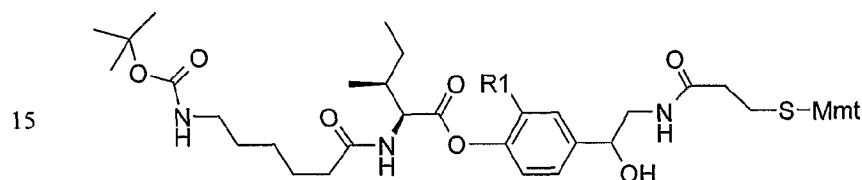
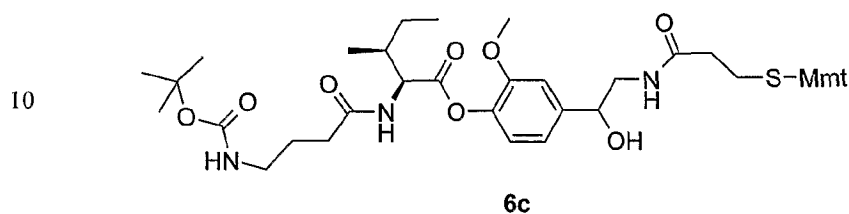
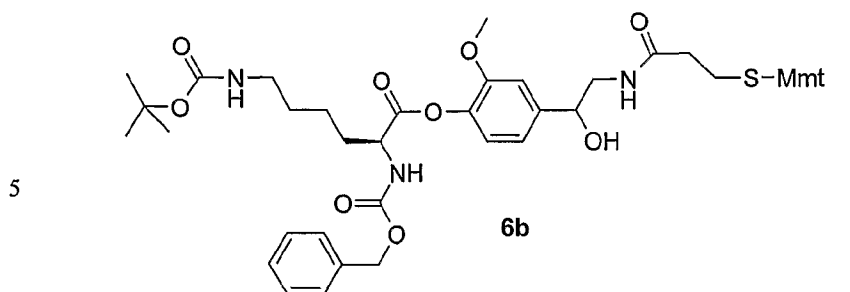
5 Synthesis of compound 6a



Mercaptothiazolide **4** (1 eq), phenol **3a** (4 eq) and DMAP (4 eq) were refluxed in DCM for 2 h under nitrogen atmosphere. After neutralization with acetic acid, the solvent was removed *in vacuo* and product **6a** was purified by RP-HPLC.

15

**6a** MS  $[M+Na]^+ = 620$  (MW+Na calculated = 620.3 g/mol)

Synthesis of compounds 6b to 6e

General synthesis protocol:      **6d** R1 = OMe  
    **6e** R1 = H

20 Carboxylic acid **5a** (1eq), phenol **3b** (1 eq), DIC (1 eq), and DMAP (2 eq) in DMF were reacted for 1h at room temperature. After addition of acetic acid (4 eq) the resulting carboxylic ester **6c** was purified by RP-HPLC.

**6d** was synthesized as described above using **5b** and **3b** as starting materials.

**6b** was synthesized as described above using Z-Lys(Boc)-OH and **3b**.

**6e** was synthesized as described above using **5b** and **3a**.

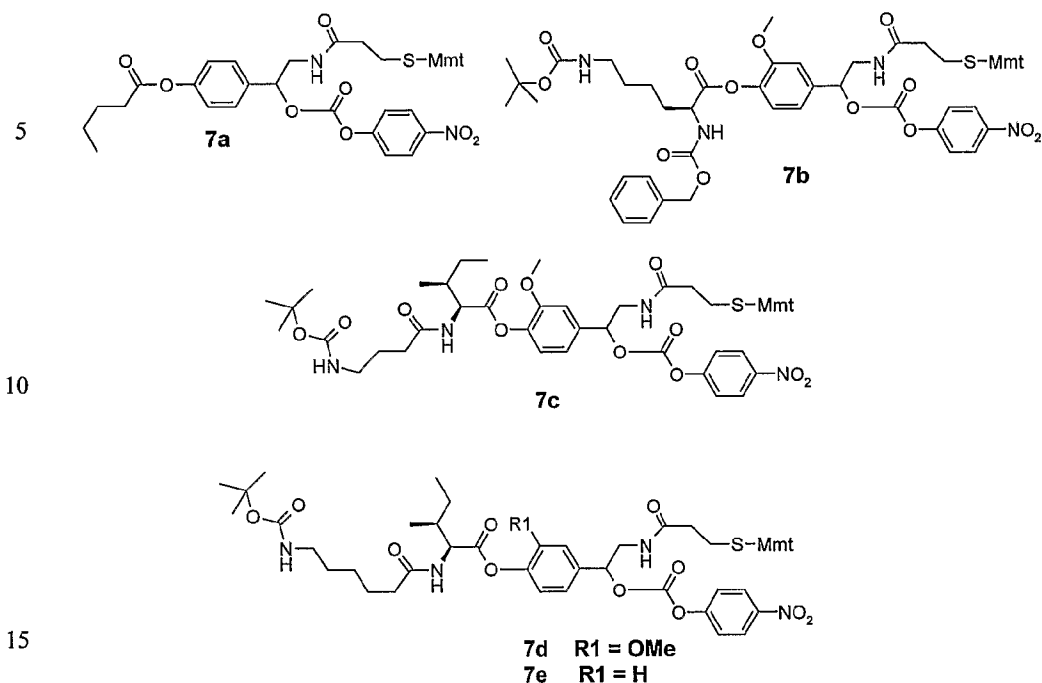
25

**6b** MS  $[M+Na]^+ = 928$  (MW+Na calculated = 928.6 g/mol)

**6c** MS  $[M+Na]^+ = 864$  (MW+Na calculated = 864.5 g/mol)

**6d** MS  $[M+Na]^+ = 892$  (MW+Na calculated = 892.6 g/mol)

30 **6e** MS  $[M+Na]^+ = 862$  (MW+Na calculated = 862.6 g/mol)

Synthesis of compounds 7a to 7e

## General synthesis protocol:

Alcohol **6a** (1 eq), 4-nitrophenyl chloroformate (10 eq), and DIEA (10 eq) were stirred in dry dioxane for 3 h at room temperature under nitrogen atmosphere. After addition of acetic acid (25 eq) the mixtures were diluted with 7/3 (v/v) acetonitrile/H<sub>2</sub>O and the carbonate **7a** was purified by RP-HPLC.

**7b**, **7c**, **7d**, or **7e** were synthesized from **6b**, **6c**, **6d**, or **6e**, respectively, as described above.

25

**7a** MS [M+Na]<sup>+</sup> = 785 (MW+Na calculated = 785.5 g/mol)

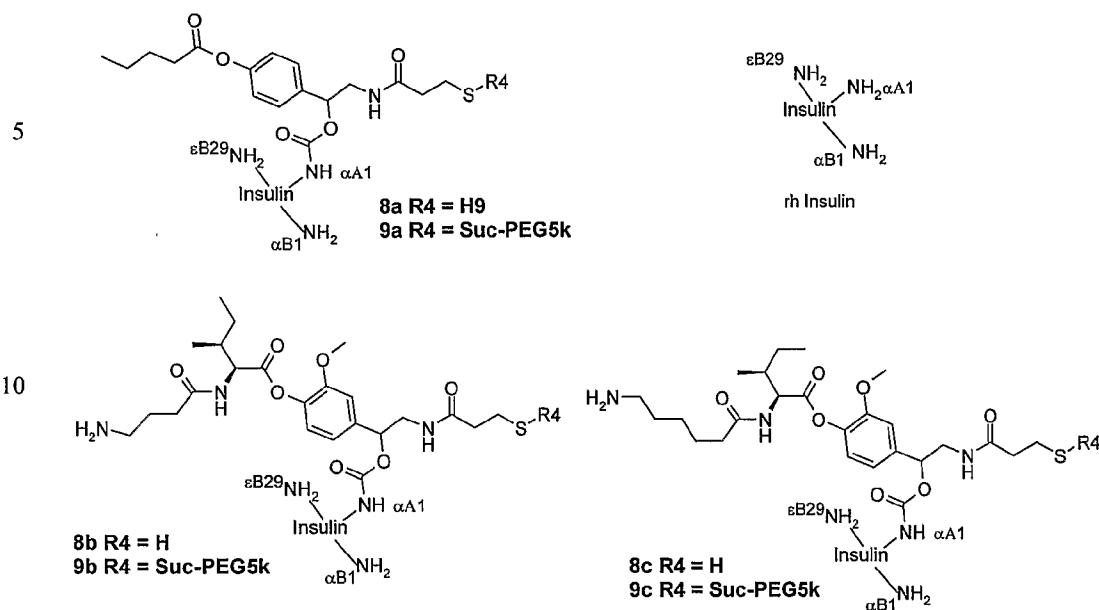
**7b** MS [M+Na]<sup>+</sup> = 1093 (MW+Na calculated = 1093.7 g/mol)

**7c** MS [M+Na]<sup>+</sup> = 1029 (MW+Na calculated = 1029.6 g/mol)

**7d** MS [M+Na]<sup>+</sup> = 1057 (MW+Na calculated = 1057.6 g/mol)

30 **7e** MS [M+Na]<sup>+</sup> = 1027 (MW+Na calculated = 1027.6 g/mol)

# Synthesis of compounds **8a** to **8c** ( $N^{\alpha A1}$ -linker-insulin)



15

General synthesis protocol:

Rh-Insulin in 1/1 (v/v) DMSO/DMF was mixed with a solution of 0.9 eq carbonate **7a** in DMSO. The resulting solution was adjusted to basic pH with DIEA and stirred for 1.5 h at RT. RP-HPLC purification gave Mmt-protected intermediate.

20 After lyophilization, the Mmt-protected intermediate was mixed with 95:5 (v/v) TFA/triethylsilane and stirred for 5 min. Volatiles were removed under nitrogen flow and **8a** was purified by RP-HPLC and lyophilized. Regioselectivity of insulin modification was verified by DTT reduction and MS analysis.

**8b** or **8c** were synthesized from **7c**, or **7d**, respectively, as described above.

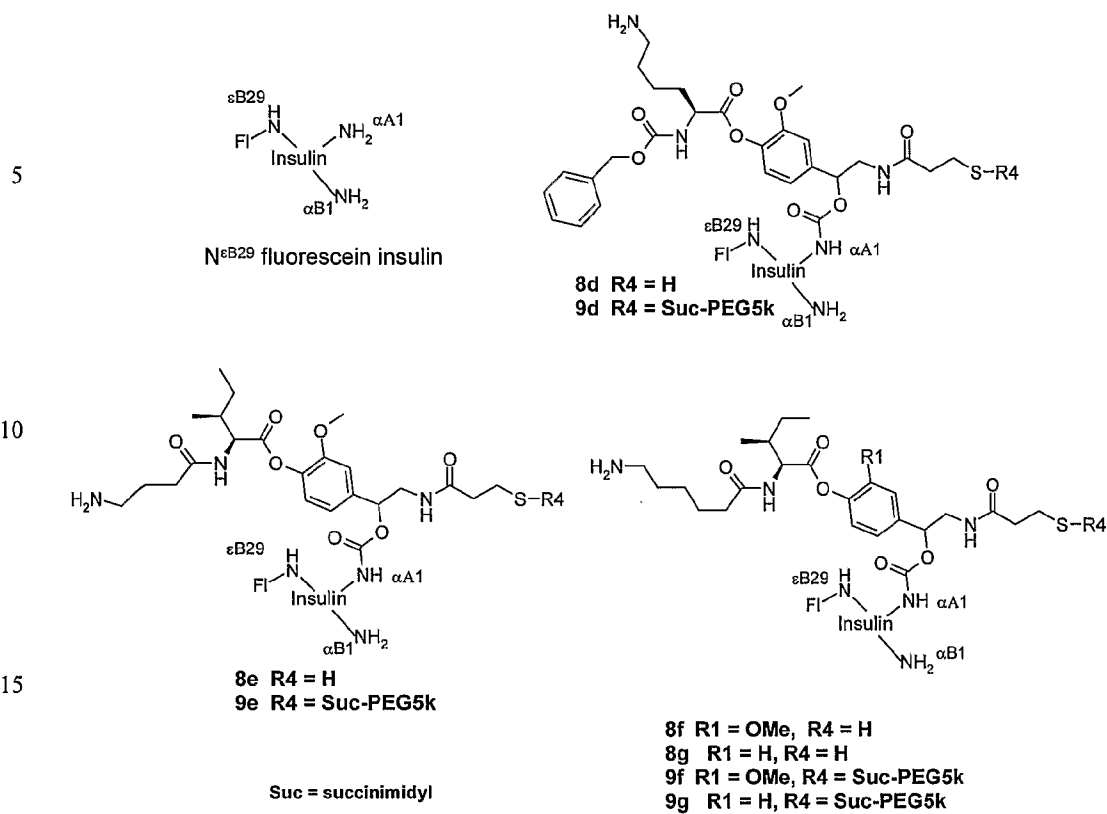
25

**8a** MS  $[M+2H]^{2+} = 3078.9$ ;  $[M+3H]^{3+} = 2053.2$   $[M+4H]^{4+} = 1540.6$  (MW calculated = 6158 g/mol)

**8b** MS  $[M+2H]^{2+} = 3152.9$ ;  $[M+3H]^{3+} = 2100.6$   $[M+4H]^{4+} = 1575.8$  (MW calculated = 6302 g/mol)

30 **8c** MS:  $[M+3H]^{3+} = 2110.7$ ;  $[M+4H]^{4+} = 1583.7$ ;  $[M+5H]^{5+} = 1266.6$  (MW calculated = 6330 g/mol)

Synthesis of compounds 8d to 8g (N<sup>εB29</sup>-Fluorescein-N<sup>αA1</sup>-linker-insulin)



Synthesis of N<sup>εB29</sup>-fluorescein insulin:

80 mg (13.8 μmol) rh-insulin were dissolved in 4 ml 1/1 (v/v) DMF/DMSO and 40 μl DIEA were added. 8 mg (17 μmol) 5-(and-6)-carboxyfluorescein succinimidyl ester were added and the solution was stirred for 30 min at room temperature. 4 ml 5/5/1 (v/v/v) acetonitrile/water/acetic acid were added, product N<sup>εB29</sup>-fluorescein insulin was purified by RP-HPLC and lyophilized. The conjugation site was verified by reduction of N<sup>εB29</sup>-fluorescein insulin with 1,4-dithiothreitol, protease digestion and MS analysis.

MS: [M+2H]<sup>2+</sup> = 3084.0; [M+3H]<sup>3+</sup> = 2054.6 (MW calculated = 6166 g/mol)

N<sup>ε</sup>B<sup>29</sup>-fluorescein insulin in 1/1 (v/v) DMF/DMSO was mixed with a solution of 0.9 eq carbonate **7b** in DMSO. The resulting solution was adjusted to basic pH with DIEA and stirred for 3 h at RT. RP-HPLC purification gave Mmt-protected intermediate.

- 5 After lyophilization, the intermediate was dissolved in 95/5 (v/v) TFA/triethylsilane and stirred for 5 min. Volatiles were removed under nitrogen flow and **8d** was purified by RP-HPLC and lyophilized.

**8e**, **8f**, or **8g** were synthesized as described above using **7c**, **7d**, or **7e**, respectively.

- 10 **8d** MS:  $[M+2H]^{2+} = 3364.1$ ;  $[M+3H]^{3+} = 2242.7$ ;  $[M+4H]^{4+} = 1681.5$  (MW calculated = 6724 g/mol)

**8e** MS:  $[M+3H]^{3+} = 2219.2$   $[M+4H]^{4+} = 1665.9$ ;  $[M+5H]^{5+} = 1332.8$  (MW calculated = 6660 g/mol)

- 8f** MS:  $[M+3H]^{3+} = 2229.7$   $[M+4H]^{4+} = 1673.3$ ;  $[M+5H]^{5+} = 1337.7$  (MW calculated = 6689 g/mol)

**8g** MS:  $[M+3H]^{3+} = 2218.7$   $[M+4H]^{4+} = 1664.9$  (MW calculated = 6659 g/mol)

#### Synthesis of compounds **9a** to **9g** (mono-pegylated insulin compounds)

20

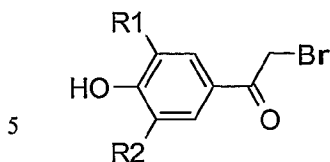
70  $\mu$ l 500  $\mu$ M **8a** in 1/4 (v/v) acetonitrile/water were mixed with 7  $\mu$ l 10 mM maleimide-PEG5k in 1/4 (v/v) acetonitrile/water and 10  $\mu$ l 0.5 M sodium phosphate buffer pH 7.0 and incubated for 15 min. Compound **9a** was purified by SEC (column: Superdex 200, flow rate: 0.75 ml/min) using 10 mM HEPES buffer (pH 7.4), 150 mM

25 NaCl, 3 mM EDTA, and 0.005% Tween as mobile phase. The collected eluate (approximately 1.5 ml) was directly used as such for release rate determination.

**9b**, **9c**, **9d**, **9e**, **9f**, or **9g** were synthesized as described above from **8b**, **8c**, **8d**, **8e**, **8f**, or **8g**, respectively.

- 30 **9a** through **9g**: SEC retention time: 19.5 min



Synthesis of compounds **11a** and **11b**

**11a** R1 = R2 = Me

**11b** R1 = Me, R2 = H

- 10 3,5-Dimethyl-4-hydroxy-acetophenone (5.0 mmol) (**10a**) and CuBr<sub>2</sub> (1.7 g, 7.5 mmol) were dissolved in 10 ml ethyl acetate and refluxed for 2 h. Solid byproducts were removed by filtration. The filtrate was evaporated and crude product **11a** was purified by RP-HPLC.

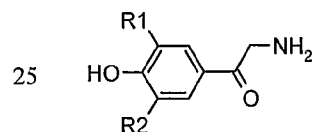
- 11b** was synthesized from 4-hydroxy-3-methyl-acetophenone (**10b**) (0.75 g, 5.0 mmol) as described above.
- 15

**11a**: Yield 754 mg (62%)

MS [M+H]<sup>+</sup> = 243.1/245.1 (MW+H calculated = 244.1 g/mol)

**11b**: Yield 533 mg (47%)

- 20 MS [M+H]<sup>+</sup> = 229.2/231.1 (MW+H calculated = 230.1 g/mol)

Synthesis of compounds **12a** and **12b**

**12a** R1 = R2 = Me

**12b** R1 = Me, R2 = H

- 30 500 mg **11a** (2.06 mmol) and 576 mg (4.11 mmol) hexamethylenetetramine were dissolved in 20 ml of trichloromethane and refluxed for 30 min. The solvent was removed in vacuo. 4 ml ethanol and 2 ml of concentrated HCl were added and the slurry was heated to 50 °C for 4 h. The mixture was concentrated in vacuo, diluted with acetonitrile/water and **12a** was purified by RP-HPLC.

**12b** was synthesized from 472 mg (2.06 mmol) **11b** as described above.

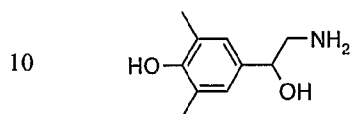
**12a**: Yield 547 mg (81%) as TFA-salt

MS  $[M+Na]^+ = 202.2$  (MW+Na calculated = 202.2 g/mol)

5 **12b**: Yield 455 mg (70%) as TFA-salt

MS  $[M+Na]^+ = 188.2$  (MW+Na calculated = 188.2 g/mol)

### Synthesis of compound 13



**13**

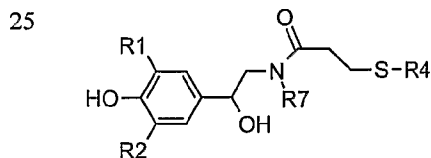
500 mg (1.71 mmol) **12a** (TFA salt) were dissolved in 10 ml 1/1 (v/v)  
15 methanol/water, 129 mg (3.41 mmol) NaBH<sub>4</sub> were added and the mixture was stirred  
for 30 min at RT. 0.5 ml of acetic acid were added and **13** was purified by RP-HPLC.

**13**: Yield 313 mg (62%) as TFA-salt

MS  $[M+Na]^+ = 204.2$  (MW+Na calculated = 204.2 g/mol)

20 NMR (300 MHz, DMSO-d<sub>6</sub>)  $\delta$ [ppm] = 8.25 (s, 1H, Phenol), 7.84 (bs, 3H, NH<sub>3</sub><sup>+</sup>),  
6.89 (s, 2H, CH<sub>ar</sub>), 5.85 (d, 1H, Hydroxyl, J=3.7 Hz), 4.62 (m, 1H, CH<sub>Benzyl</sub>), 2.93 (m,  
1H, CH<sub>a</sub>), 2.80 (m, 1H, CH<sub>b</sub>), 2.17 (s, 6H, CH<sub>3</sub>).

### Synthesis of compounds 14a to 14d



**14a** R1 = R2 = Me, R7 = H, R4 = Mmt

30 **14b** R1 = R2 = H, R7 = Me, R4 = Mmt

**14c** R1 = OMe, R2 = H, R7 = Me, R4 = Mmt

**14d** R1 = H, R2 = H, R7 = Me, R4 = Trt

**13** (TFA salt, 159 mg, 0.541 mmol) was coupled to compound **2** as described for compound **3a** to yield **14a**.

**14b**, or **14c** were synthesized as described above using synephrine (335 mg, 2.00 mmol) or metanephrine (HCl salt, 281 mg, 1.20 mmol), respectively.

- 5 Synephrine (335 mg, 2.3 mmol) was coupled to 3-tritylsulfanyl-propionic acid as described above to yield **14d**.

**14a**: Yield 254 mg (87%)

MS  $[M+Na]^+ = 564.7$  (MW+Na calculated = 564.3 g/mol)

- 10 **14b**: Yield 760 mg (72%)

MS  $[M+Na]^+ = 550.2$  (MW+Na calculated = 550.3 g/mol)

**14c**: Yield 530 mg (80%)

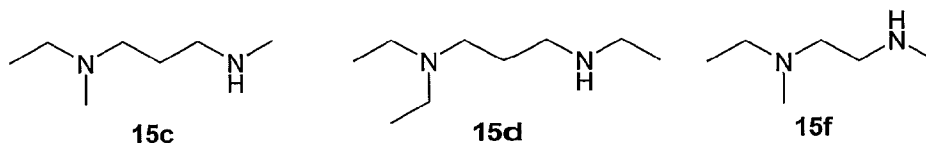
MS  $[M+Na]^+ = 580.4$  (MW+Na calculated = 580.4 g/mol)

**14d**: Yield 567 mg (49%)

- 15 MS  $[M+Na]^+ = 520.5$  (MW+Na calculated = 520.7 g/mol)

#### Synthesis of compounds **15c**, **15d** and **15f**

20



General synthesis protocol:

- 25 1 g 2-chlorotrityl chloride resin (loading 1.4 mmol/g) was incubated for 1 h with N,N'-dimethylpropane-1,3-diamine (for synthesis of **15c**), or N,N'-diethylpropane-1,3-diamine (for synthesis of **15d**) or N,N'-dimethylethane-1,2-diamine (4 eq) (for synthesis of **15f**) in DCM. After washing of the resin with DMF, amines were acetylated with 1/1/2 (v/v/v) acetic anhydride/pyridine/DMF for 14 h. The resin was washed with THF and dried. LiAlH<sub>4</sub> (1 M in THF, 4 eq) was added dropwise to the suspended resin in THF. The resulting suspension was stirred for 3 h at 45°C under nitrogen atmosphere. After cooling, aqueous Rochelle's salt solution was added and the resin was separated and dried. Compounds were cleaved from resin with 2/1 (v/v)
- 30

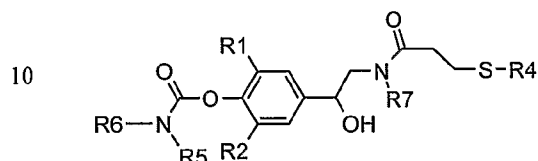
HFIP/DCM (2 x 30 min). The volatile components were evaporated and the products **15c**, **15d** or **15f** were used in the following steps without further purification.

**15c** MS  $[M+H]^+ = 131.2$  (MW = 130.1 g/mol)

5 **15d** MS  $[M+H]^+ = 159.2$  (MW = 158.1 g/mol)

**15f** MS  $[M+H]^+ = 117.1$  (MW = 116 g/mol)

#### Synthesis of compounds **16a** to **16f** and **16i**



**16a** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl, R7 = H, R4 = Mmt

**16b** R1 = OMe, R2 = H, R5 = Et, R6 = 2-(diethylamino)ethyl, R7 = H, R4 = Mmt

15 **16c** R1 = OMe, R2 = H, R5 = Me, R6 = 3-(N-ethyl-N-methylamino)propyl, R7 = Me, R4 = Mmt

**16d** R1 = R2 = H, R5 = Me, R6 = 3-(N-ethyl-N-methylamino)propyl, R7 = Me, R4 = Mmt

**16e** R1 = OMe, R2 = H, R5 = Et, R6 = 3-(diethylamino)propyl, R7 = Me, R4 = Mmt

20 **16f** R1 = R2 = H, R5 = Et, R6 = 3-(diethylamino)propyl, R7 = Me, R4 = Mmt

**16i** R1 = R2 = H, R5 = Et, R6 = 2-(diethylamino)ethyl, R7 = Me, R4 = Trt

**14a** (120 mg, 0.222 mmol) was dissolved in 1.5 ml of dry THF. p-Nitrophenyl-chloroformate (45 mg, 0.222 mmol) and DIEA (113  $\mu$ l, 0.665 mmol) were added and the mixture was stirred for 30 min at RT. **15a** (N,N,N'-trimethyl-ethylene-1,2-diamine) (72  $\mu$ l, 0.554 mmol) was added and stirring was continued for 30 min. The solvent was removed in vacuo, 100  $\mu$ l of AcOH were added and **16a** was purified by RP-HPLC.

30 **16b** was synthesized as described above from **3b** (80 mg, 0.15 mmol) and **15b** (N,N,N'-triethyl-ethylene-1,2-diamine) (55 mg, 0.38 mmol).

**16c** or **16d** were synthesized as describe above from **14c** (56 mg, 0.1 mmol) or **14b** (53 mg, 0.1 mmol), respectively, and diamine **15c**.

**16e** or **16f** were synthesized as described above from **14c** (56 mg, 0.1 mmol) or **14b**, respectively, (53 mg, 0.1 mmol) and diamine **15d**.

**16i** was synthesized as described above from **14d** (350 mg, 0.7 mmol) and **15b** (N,N,N'-triethyl-ethylene-1,2-diamine) (180  $\mu$ l, 1 mmol).

5

**16a:** Yield 120 mg (69%) as TFA salt

MS  $[M+Na]^+ = 692.4$  (MW+Na calculated = 692.9 g/mol)

**16b:** Yield 48 mg (40%) as TFA salt

MS  $[M+Na]^+ = 736.3$  (MW+Na calculated = 736.4 g/mol)

10 

**16c:** Yield 8 mg (10%) as TFA salt

MS  $[M+Na]^+ = 736.4$  (MW+Na calculated = 736.4 g/mol)

**16d:** Yield 20 mg (25%) as TFA salt

MS  $[M+Na]^+ = 706.3$  (MW+Na calculated = 706.3 g/mol)

**16e:** Yield 2 mg (3%) as TFA salt

15 

MS  $[M+Na]^+ = 764.6$  (MW+Na calculated = 764.4 g/mol)

**16f:** Yield 6 mg (8%) as TFA salt

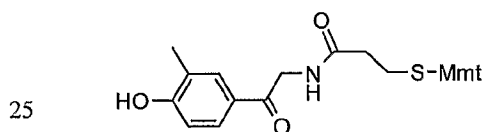
MS  $[M+Na]^+ = 734.4$  (MW+Na calculated = 734.3 g/mol)

**16i:** Yield 152 mg (28%) as TFA salt

MS  $[M+Na]^+ = 690.5$  (MW+Na calculated = 690.9 g/mol)

20

### Synthesis of compound 17



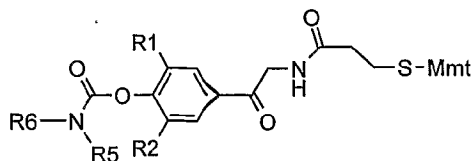
**17**

Amine **12b** (TFA salt) was coupled to compound **2** as described for compound **3a**.

30

**17:** Yield 608 mg (74%)

MS  $[M+Na]^+ = 548.3$  (MW+Na calculated = 548.7 g/mol)

Synthesis of compounds 18a and 18b

**18a** R1 = R5 = Me, R2 = H, R6 = 3-(dimethylamino)propyl

**18b** R1 = R5 = Me, R2 = H, R6 = 2-(N-ethyl-N-methylamino)ethyl

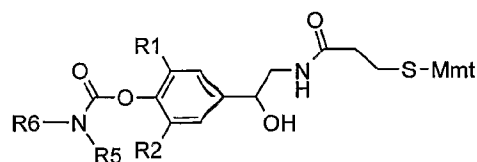
383 mg (0.729 mmol) of **17** were reacted with p-nitrophenyl-chloroformate and N,N,N'-trimethyl-propane-1,3-diamine (**15e**) or **15f**, respectively, to yield **18a** or **18b** as described for compound **16a**.

**18a**: Yield 287 mg (50%) as TFA salt

MS  $[M+Na]^+ = 690.7$  (MW+Na calculated = 690,9 g/mol)

**18b**: Yield 148 mg (26%) as TFA salt

MS  $[M+Na]^+ = 690.9$  (MW+Na calculated = 690,9 g/mol)

Synthesis of compounds 16g and 16h

**16g** R1 = R5 = Me, R2 = H, R6 = 3-(dimethylamino)propyl

**16h** R1 = R5 = Me, R2 = H, R6 = 2-(N-ethyl-N-methylamino)ethyl

**18a** (287 mg, 0.367 mmol, TFA salt) was dissolved in 5 ml methanol, NaBH<sub>4</sub> (41 mg, 1.07 mmol) was added and the mixture was stirred for 30 min at RT. 0.5 ml of acetic acid were added and **16g** was purified by RP-HPLC.

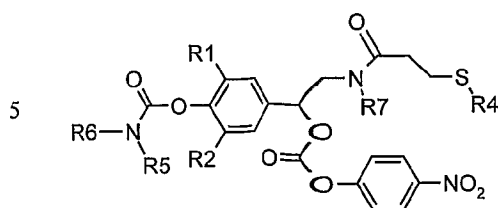
**18b** (8 mg, 0.010 mmol, TFA salt) was reacted as described above to yield **16h**.

**16g**: Yield 201 mg (70%) as TFA-salt

MS  $[M+Na]^+ = 692.7$  (MW+Na calculated = 692.9 g/mol)

**16h**: Yield 6 mg (77%) as TFA-salt

MS  $[M+Na]^+ = 692.7$  (MW+Na calculated = 692.9 g/mol)

Synthesis of compounds 19a to 19i

**19a** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl, R7 = H, R4 = Mmt

**19b** R1 = OMe, R2 = H, R5 = Et, R6 = 2-(diethylamino)ethyl, R7 = H, R4 = Mmt

10 **19c** R1 = OMe, R2 = H, R5 = Me, R6 = 3-(N-ethyl-N-methylamino)propyl, R7 = Me, R4 = Mmt

**19d** R1 = R2 = H, R5 = Me, R6 = 3-(N-ethyl-N-methylamino)propyl, R7 = Me, R4 = Mmt

**19e** R1 = OMe, R2 = H, R5 = Et, R6 = 3-(diethylamino)propyl, R7 = Me, R4 = Mmt

15 **19f** R1 = R2 = H, R5 = Et, R6 = 3-(diethylamino)propyl, R7 = Me, R4 = Mmt

**19g** R1 = R5 = Me, R2 = H, R6 = 3-(dimethylamino)propyl, R7 = H, R4 = Mmt

**19h** R1 = R5 = Me, R2 = H, R6 = 2-(N-ethyl-N-methylamino)ethyl, R7 = H, R4 = Mmt

**19i** R1 = R2 = H, R5 = Et, R6 = 2-(diethylamino)ethyl, R7 = Me, R4 = Trt

20

Carbonates **19a** to **19i** were synthesized from **16a** to **16i**, respectively, as described for compound **7a**.

**19a:** Yield 98 mg (72%) as TFA-salt

25 MS [M+Na]<sup>+</sup> = 857.8 (MW+Na calculated = 858.0 g/mol)

**19b:** Yield 6 mg (11 %) as TFA-salt

MS [M+Na]<sup>+</sup> = 901.8 (MW+Na calculated = 901.5 g/mol)

**19c:** Yield 1 mg (15 %) as TFA-salt

MS [M+Na]<sup>+</sup> = 901.4 (MW+Na calculated = 901.5 g/mol)

30 **19d:** Yield 8 mg (29%) as TFA-salt

MS [M+Na]<sup>+</sup> = 871.4 (MW+Na calculated = 871.4 g/mol)

**19e:** Yield 0.3 mg (18%) as TFA-salt

MS [M+Na]<sup>+</sup> = 929.4 (MW+Na calculated = 929.5 g/mol)

**19f:** Yield 4 mg (45%) as TFA-salt

MS  $[M+Na]^+ = 899.7$  (MW+Na calculated = 899.6 g/mol)

**19g:** Yield 6 mg (6%) as TFA-salt

MS  $[M+Na]^+ = 857.8$  (MW+Na calculated = 858.0 g/mol)

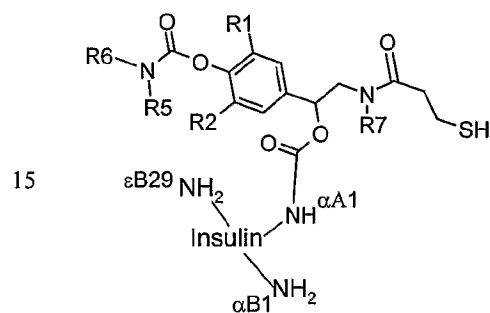
**19h:** Yield 0.8 mg (11%) as TFA-salt

5 MS  $[M+Na]^+ = 857.7$  (MW+Na calculated = 858.0 g/mol)

**19i:** Yield 77 mg (49%) as TFA-salt

MS  $[M+Na]^+ = 856.2$  (MW+Na calculated = 856.0 g/mol)

10 Synthesis of compounds **20a** to **20f**



**20a** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl, R7 = H

20 **20b** R1 = OMe, R2 = H, R5 = Et, R6 = 2-(diethylamino)ethyl, R7 = H

**20c** R1 = OMe, R2 = H, R5 = Me, R6 = 3-(N-ethyl-N-methylamino)propyl, R7 = Me

**20d** R1 = R2 = H, R5 = Me, R6 = 3-(N-ethyl-N-methylamino)propyl, R7 = Me

**20e** R1 = R5 = Me, R2 = H, R6 = 3-(dimethylamino)propyl, R7 = H

**20f** R1 = R5 = Me, R2 = H, R6 = 2-(N-ethyl-N-methylamino)ethyl, R7 = H

25

Insulin derivatives **20a**, **20b**, **20c**, **20d**, **20e**, or **20f** were synthesized from **19a**, **19b**, **19c**, **19d**, **19g**, or **19h** respectively, as described for compound **8a**.

**20a** MS  $[M+3H]^{3+} = 2077.3$   $[M+4H]^{4+} = 1559.2$  (MW calculated = 6231.3 g/mol)

30 **20b** MS  $[M+3H]^{3+} = 2093.0$   $[M+4H]^{4+} = 1569.6$  (MW calculated = 6274 g/mol)

**20c** MS  $[M+3H]^{3+} = 2090.8$   $[M+4H]^{4+} = 1568.7$  (MW calculated = 6274 g/mol)

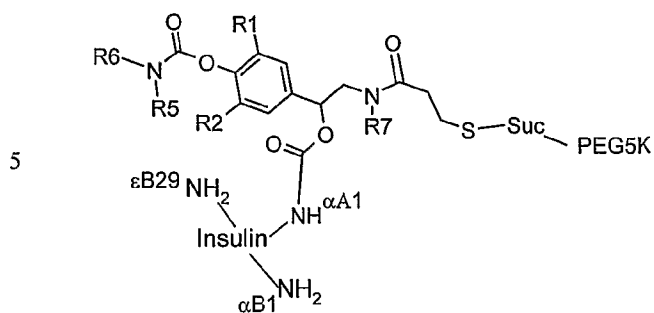
**20d** MS  $[M+3H]^{3+} = 2081.3$   $[M+4H]^{4+} = 1561.8$  (MW calculated = 6244 g/mol)

**20e** MS  $[M+3H]^{3+} = 2077.1$   $[M+4H]^{4+} = 1558.2$  (MW calculated = 6231.3 g/mol)

**20f** MS  $[M+3H]^{3+} = 2076.7$   $[M+4H]^{4+} = 1559.3$  (MW calculated = 6231.3 g/mol)



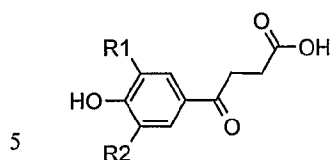
### Synthesis of compounds **21a** to **21f** (mono-pegylated insulin derivatives)



- 10    **21a** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl, R7 = H  
       **21b** R1 = OMe, R2 = H, R5 = Et, R6 = 2-(diethylamino)ethyl, R7 = H  
       **21c** R1 = OMe, R2 = H, R5 = Me, R6 = 3-(N-ethyl-N-methylamino)propyl, R7 = Me  
       **21d** R1 = R2 = H, R5 = Me, R6 = 3-(N-ethyl-N-methylamino)propyl, R7 = Me  
       **21e** R1 = R5 = Me, R2 = H, R6 = 3-(dimethylamino)propyl, R7 = H  
 15    **21f** R1 = R5 = Me, R2 = H, R6 = 2-(N-ethyl-N-methylamino)ethyl, R7 = H

Insulin derivatives **21a**, **21b**, **21c**, **21d**, **21e**, or **21f** were synthesized from compound **20a**, **20b**, **20c**, **20d**, **20e**, or **20f**, respectively, as described for compound **9a**.

- 20     **21a** through **21f**: SEC retention time: 19.5 min

Synthesis of compounds 23a and 23b

**23a** R1 = Me, R2 = H

**23b** R1 = R2 = Me

10 o-Cresol (**22a**) (1 eq), succinic anhydride (1 eq), and AlCl<sub>3</sub> (3 eq) in nitrobenzene were heated to 100 °C for 1 h. The reaction mixture was poured on HCl/ice and extracted with ether. The organic layer was extracted with 1 N NaOH and the aqueous layer was acidified with concentrated HCl. The aqueous layer was extracted with ether and the ether was evaporated. **23a** was purified by RP-HPLC.

**23b** was synthesized from 2,6-dimethylphenol (**22b**) as described above.

15

**23a:** Yield 552 mg (31%)

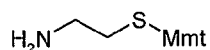
MS [M+Na]<sup>+</sup> = 231.0 (MW+Na calculated = 231.2 g/mol)

20 NMR (300 MHz, DMSO-d<sub>6</sub>) δ[ppm] = 12.05 (bs, 1H, CO<sub>2</sub>H), 10.23 (s, 1H, phenol OH), 7.74 (s, 1H, CH<sub>ar</sub>), 7.7 (d, 1H, CH<sub>ar</sub>, <sup>3</sup>J<sub>H,H</sub> = 8.4 Hz), 6.86 (d, 1H, CH<sub>ar</sub>, <sup>3</sup>J<sub>H,H</sub> = 8.4 Hz), 3.13 (t, 2H, C(O)CH<sub>2</sub>, <sup>3</sup>J<sub>H,H</sub> = 6.4 Hz), 2.53 (t, 2H, CH<sub>2</sub>CO<sub>2</sub>, <sup>3</sup>J<sub>H,H</sub> = 6.4 Hz), 2.16 (s, 3H, CH<sub>3</sub>)

**23b:** Yield 166 mg (15%)

MS [M+Na]<sup>+</sup> = 245.4 (MW+Na calculated = 245.2 g/mol)

25 Synthesis of compound 24



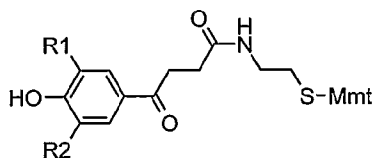
1.85 g (16.02 mmol) cysteamine hydrochloride were dissolved in 15 ml of TFA and 2.47 g (8.01 mmol) MmtCl were added. After stirring the mixture at RT for 20 min the solvent was evaporated in vacuo. The residue was dissolved in diethyl ether and extracted with saturated aqueous NaHCO<sub>3</sub>, 1N H<sub>2</sub>SO<sub>4</sub> and brine. The solvent was evaporated and **24** was purified by RP-HPLC.

**24:** Yield 1.11 g (30%) as TFA salt

TLC (AcOEt/Et<sub>3</sub>N 99/1), R<sub>f</sub> = 0.24

#### Synthesis of compounds **25a** and **25b**

5



**25a** R1 = Me, R2 = H

10 **25b** R1 = R2 = Me

**23a** (1eq), HOBt (1.1 eq) and DIC (1 eq) were dissolved in DMF and stirred at RT for 30 min. **24** (TFA salt, 1 eq) and DIEA (3 eq) were added and the solution was stirred for 60 min. Acetic acid was added and **25a** was purified by RP-HPLC.

15 **25b** was synthesized from **23b** as described above.

**25a:** Yield 552 mg (25%)

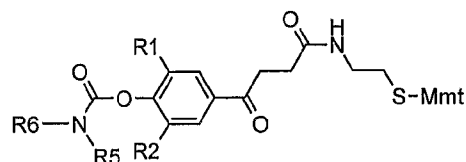
MS [M+Na]<sup>+</sup> = 562.7 (MW+Na calculated = 562.7 g/mol)

**25b:** Yield 15 mg (40%)

20 MS [M+Na]<sup>+</sup> = 576.6 (MW+Na calculated = 576.6 g/mol)

#### Synthesis of compounds **26a** and **26b**

25



**26a** R1 = Me, R2 = H, R5 = R6 = 3-(dimethylamino)propyl

**26b** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl

30 267 mg (0.495 mmol) **25a** was reacted with p-nitrophenylchloroformate and N-(3-dimethylamino-propyl)-N',N'-dimethyl-propane-1,3-diamine (**15g**) to yield **26a** as described for compound **16a**.

**26b** was synthesized as described above using 15 mg **25b** and N,N,N'-trimethyl-ethane-1,2-diamine (**15a**).

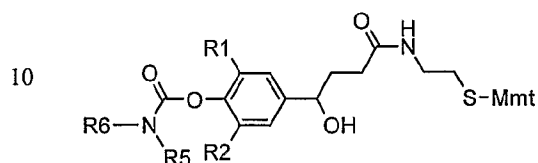
**26a:** Yield 282 mg (58%) as double TFA salt

MS  $[M+Na]^+ = 775.2$  (MW+Na calculated = 776.0 g/mol)

**26b:** Yield 17 mg (70%) as TFA salt

5 MS  $[M+Na]^+ = 704.5$  (MW+Na calculated = 704.6 g/mol)

### Synthesis of compounds 27a and 27b



**27a** R1 = Me, R2 = H, R5 = R6 = 3-(dimethylamino)propyl

**27b** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl

15

**26a** (272 mg, 0.277 mmol, double TFA salt) was dissolved in 5 ml methanol, NaBH<sub>4</sub> (42 mg, 1.09 mmol) was added and the mixture was stirred for 30 min at RT. 0.5 ml of acetic acid were added and **27a** was purified by RP-HPLC.

Alcohol **27b** was synthesized likewise from **26b** (17 mg, 25 μmol, TFA salt).

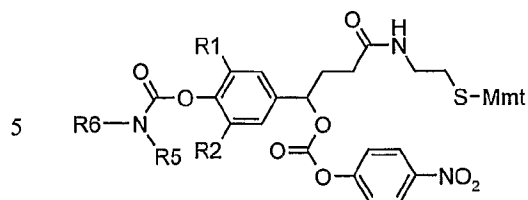
20

**27a:** Yield 142 mg (52%) as double TFA salt

MS  $[M+Na]^+ = 777.9$  (MW+Na calculated = 778.0 g/mol)

**27b:** Yield 6 mg (40%) as TFA salt

MS  $[M+Na]^+ = 706.5$  (MW+Na calculated = 706.6 g/mol)

Synthesis of compounds 28a and 28b

**28a** R1 = Me, R2 = H, R5 = R6 = 3-(dimethylamino)propyl

**28b** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl

10

Carbonates **28a** or **28b** were synthesized from **27a** or **27b**, respectively, as described for compound **7a**.

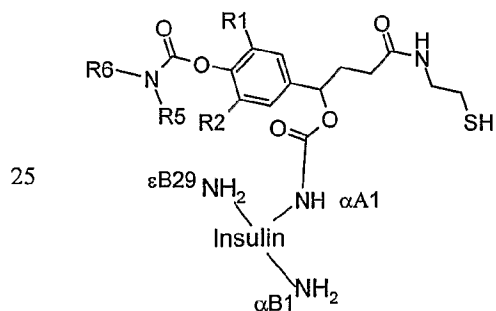
15 **28a**: Yield 1 mg (29%)

MS  $[M+Na]^+ = 942.9$  (MW+Na calculated = 943.2 g/mol)

**28b**: Yield 1.5 mg (19%)

MS  $[M+Na]^+ = 871.6$  (MW+Na calculated = 871.7 g/mol)

20 Synthesis of compounds 29a and 29b



**29a** R1 = Me, R2 = H, R5 = R6 = 3-(dimethylamino)propyl

30 **29b** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl

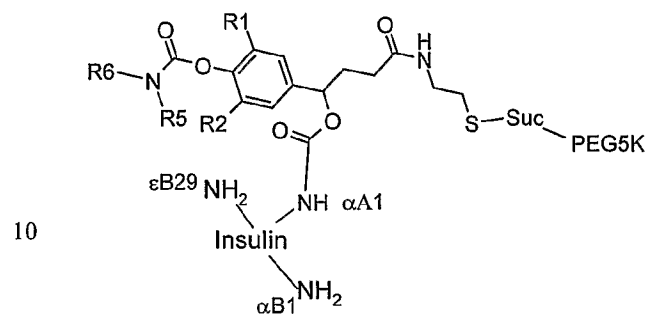
Insulin derivatives **29a** or **29b** were synthesized from **28a** or **28b**, respectively, as described for compound **8a**.

**29a** MS  $[M+3H]^{3+} = 2105.8$   $[M+4H]^{4+} = 1580.2$  (MW calculated = 6316.4 g/mol)

**29b** MS  $[M+3H]^{3+} = 2081.8$   $[M+4H]^{4+} = 1562.4$  (MW calculated = 6244 g/mol)

### Synthesis of mono-pegylated insulin derivatives **30a** and **30b**

5



10

**30a** R1 = Me, R2 = H, R5 = R6 = 3-(dimethylamino)propyl

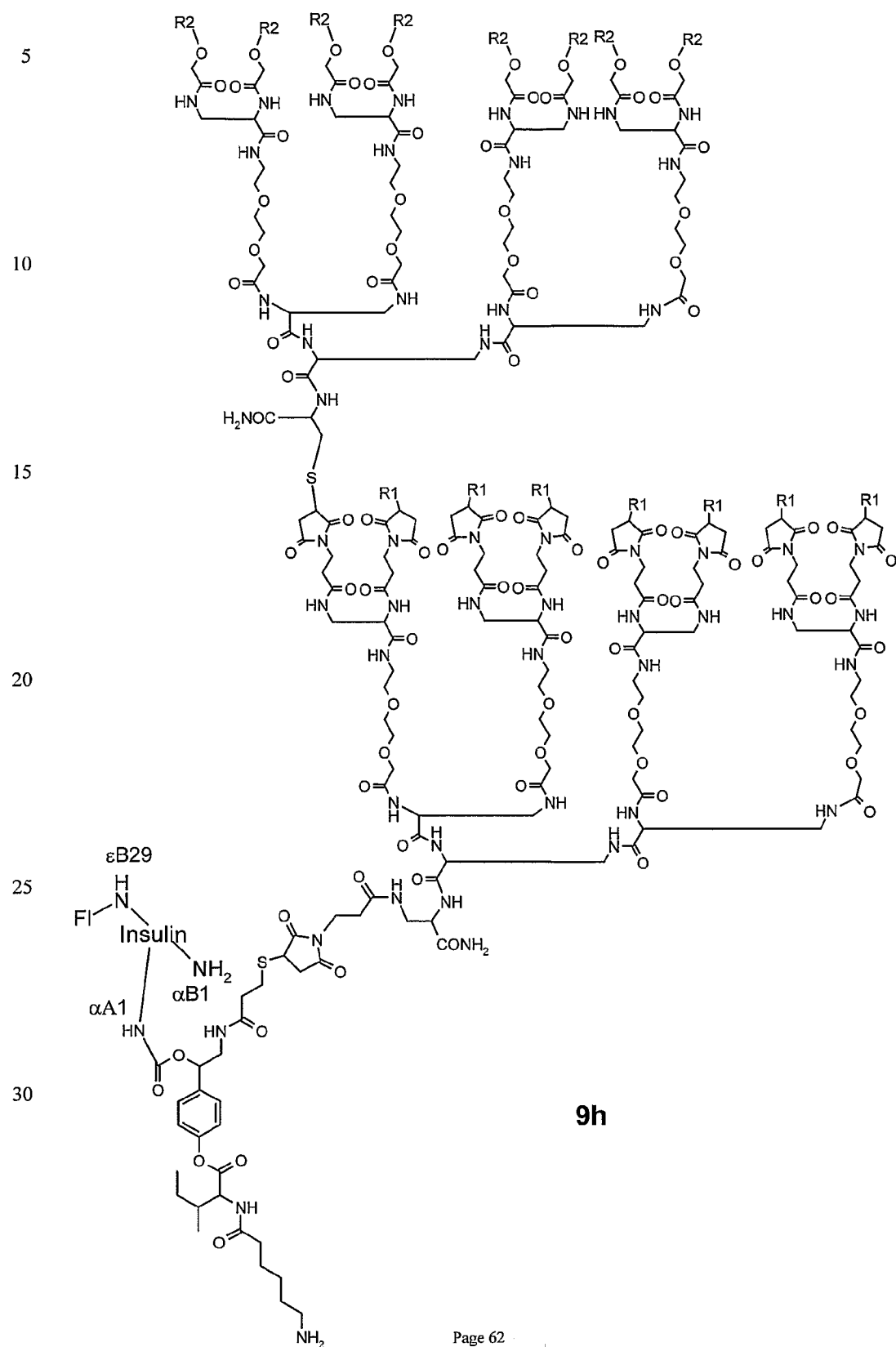
**30b** R1 = R2 = R5 = Me, R6 = 2-(dimethylamino)ethyl

15

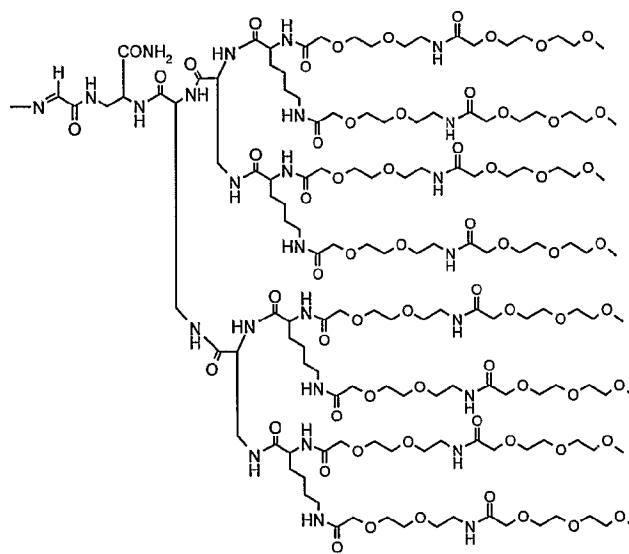
Insulin derivatives **30a** or **30b** were synthesized from **29a** or **29b**, respectively, as described for compound **9a**.

20 **30a** and **30b**: SEC retention time: 19.5 min

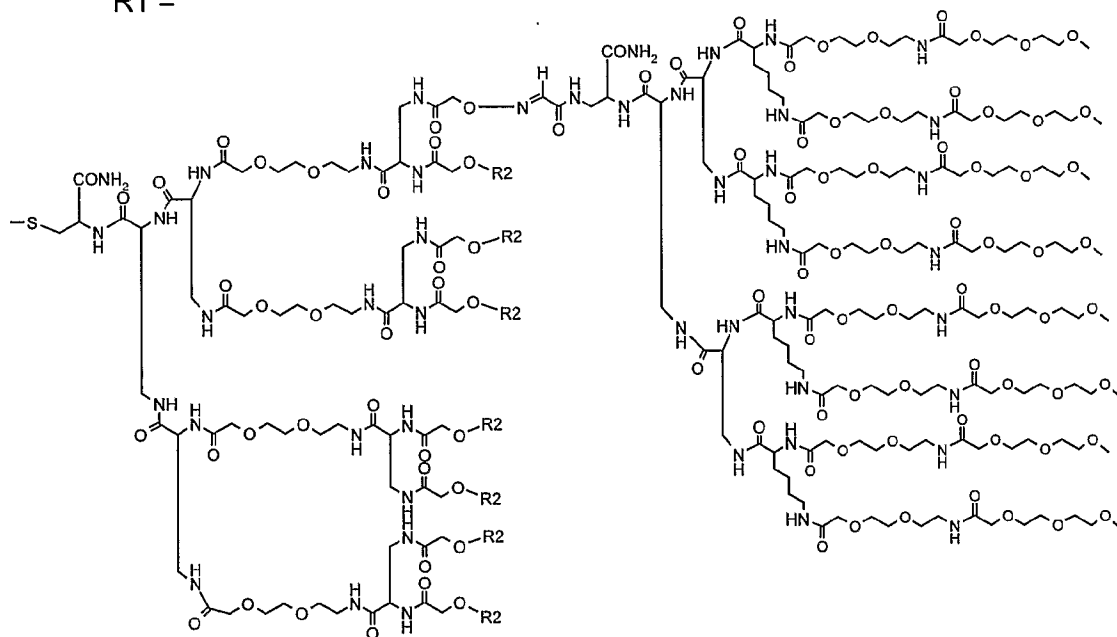
### Synthesis of a polymeric prodrug according to formula Ia with an ester-linked masking group and a dendritic carrier (9h)



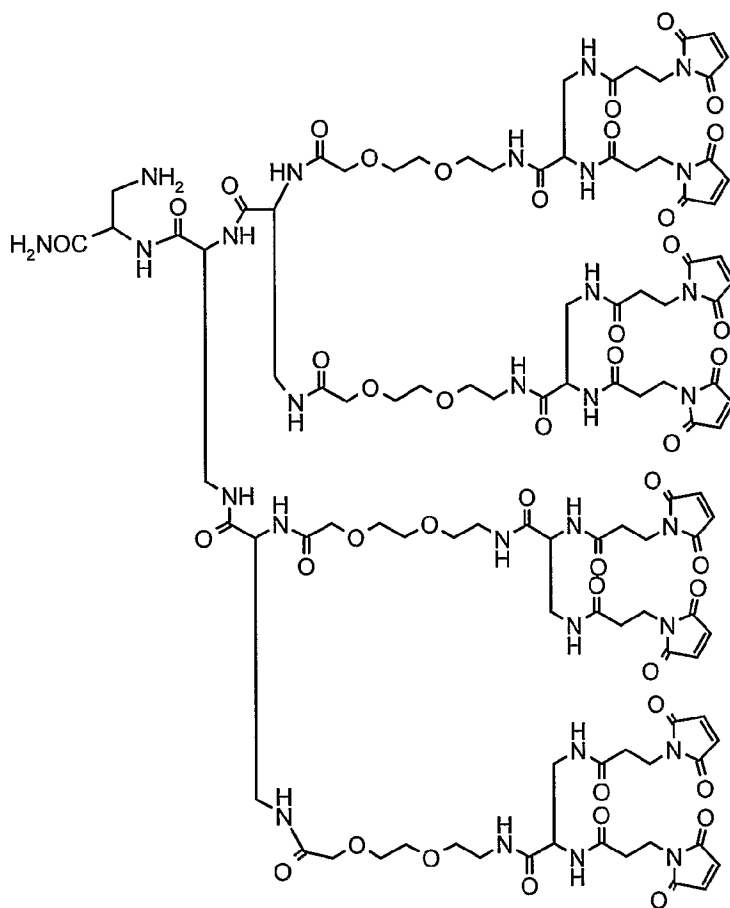
R2 =



R1 =





Synthesis of compound **31****31**

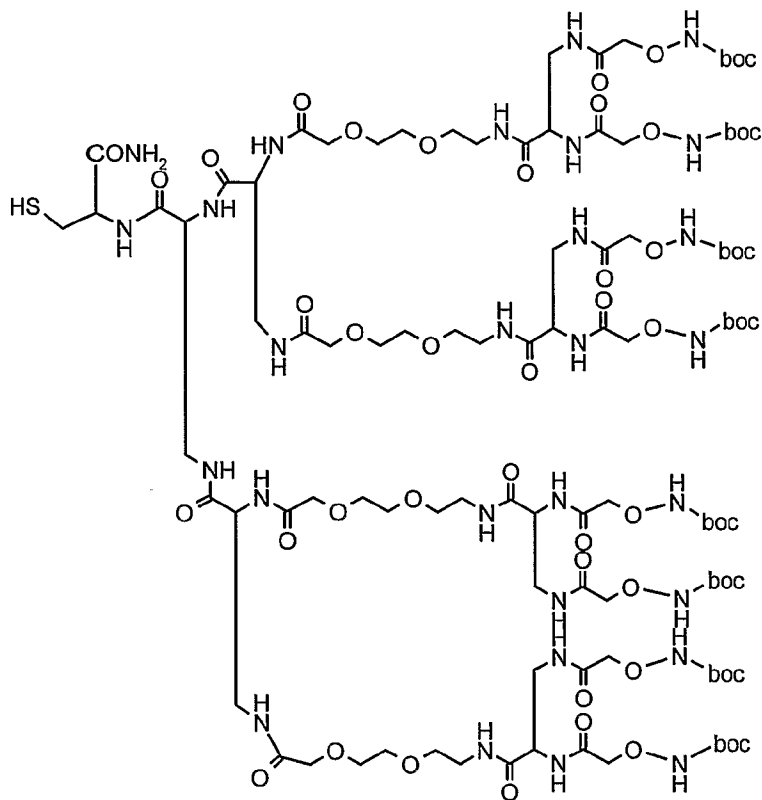
5

**31** was obtained according to the standard solid-phase synthesis protocol. The amino acids Fmoc-Dpr(Boc)-OH, Fmoc-Dpr(Fmoc)-OH, Fmoc-Dpr(Fmoc)-OH, Fmoc-Ado-OH, and Fmoc-Dpr(Fmoc)-OH were coupled to NovaSyn TG Sieber amide resin.

- 10 After final fmoc removal the resin was agitated with 5 eq maleimidopropionic acid and 5 eq DIC in relation to amino groups in DMF for 30 min. **31** was cleaved from resin with TFA/TES/water 95/3/2 (v/v/v). After evaporation of solvent, product **31** was purified by RP-HPLC.

MS:  $[M+H]^+ = 2494.6$  (MW calculated = 2495.4 g/mol)

### Synthesis of compound **32**



### **32**

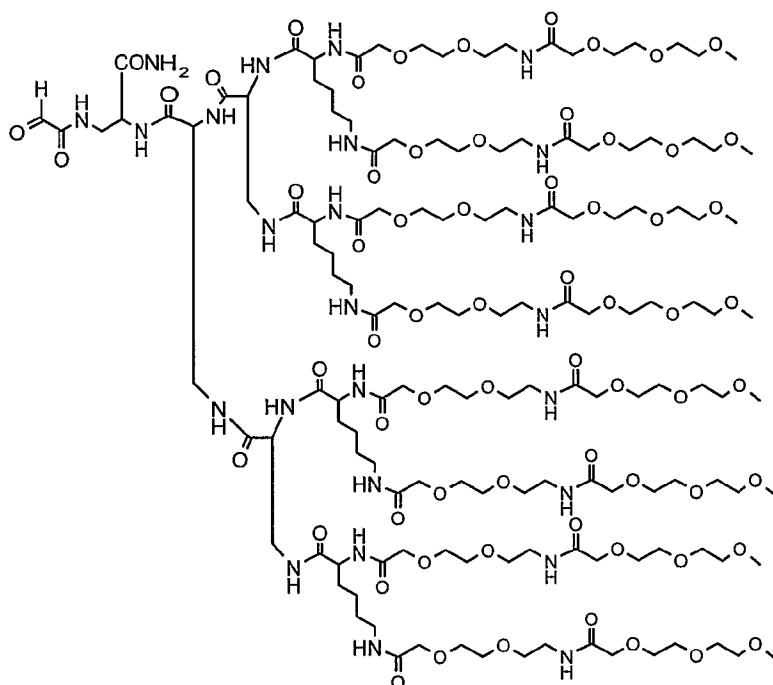
Compound **32** was obtained according to the standard solid-phase synthesis protocol.

- 5 The amino acids Fmoc-Cys(Mmt)-OH, Fmoc-Dpr(Fmoc)-OH, Fmoc-Dpr(Fmoc)-OH, Fmoc-Ado-OH, and Fmoc-Dpr(Fmoc)-OH were coupled to NovaSynTG Sieber amide resin.

- After final fmoc removal the resin was agitated with 3 eq Boc-aminoxyacetic acid, 3 eq DIC, and 3 eq HOBt in relation to amino groups in DMF for 30 min. **32** was
- 10 cleaved from resin with DCM/TFA/TES 97/1/2 (v/v/v). After addition of 0.8 eq pyridine in relation to TFA the solvent was evaporated and product **32** was purified by RP-HPLC.

MS:  $[M+H]^+ = 2688.2$  g/mol (MW calculated = 2688.8 g/mol)

## Synthesis of compound 33

**33**

5

Compound **33** was obtained according to the standard solid-phase synthesis protocol. The amino acids Fmoc-Dpr(ivDde)-OH, Fmoc-Dpr(Fmoc)-OH, Fmoc-Dpr(Fmoc)-OH, Fmoc-Lys(Fmoc)-OH, and Fmoc-Ado-OH were coupled to NovaSyn TG Sieber amide resin.

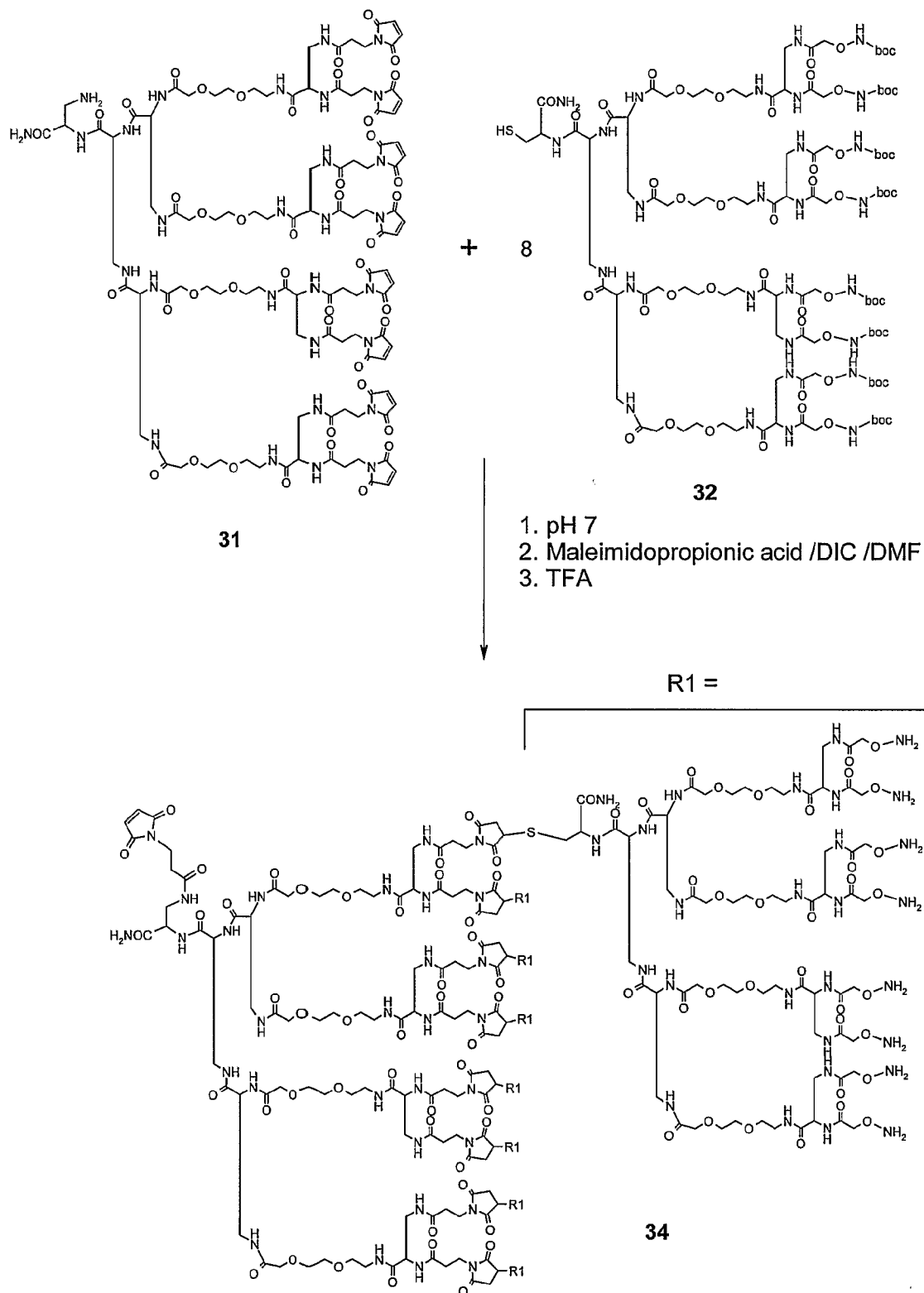
10 After final fmoc removal the resin was agitated with 3 eq 3,6,9-trioxadecanoic acid, 3 eq PyBOP, and 6 eq DIEA in relation to amino groups in DMF for 60 min.

To cleave the ivDde protecting group, the resin was treated three times with 2% hydrazine in DMF. After washing, 3 eq Fmoc-Ser-OH was coupled with 3eq DIC and 3 eq HOBt for 30 min. After final fmoc removal resin was washed and the product

15 was cleaved from resin with DCM/TFA/TES 88/10/2 (v/v/v). Solvent was evaporated and the residue was oxidized with 10 eq sodium periodate in 3/2 (v/v) 0.1 M sodium phosphate pH 7/acetonitrile for 15 min to yield **33**. Product **33** was purified by RP-HPLC and lyophilized.

MS:  $[M+H]^+ = 3372.1$  g/mol (3372.8 g/mol)

Synthesis of compound **34**



6 mg (2.4  $\mu\text{mol}$ ) of compound **31** were dissolved in 1 ml 2/1 (v/v) acetonitrile/0.1 M sodium phosphate buffer pH 7 and 65 mg (24.2  $\mu\text{mol}$ ) of compound **32** were added.

5 The solution was stirred at room temperature for 2 h and then the product purified by RP-HPLC and lyophilized (yield: 45 mg (78%)).

The lyophilized product (45 mg) was dissolved in 0.5 ml DMF and 10  $\mu\text{l}$  DIEA were added. 5 mg (30  $\mu\text{mol}$ ) 3-maleimidopropionic acid and 4.7  $\mu\text{l}$  (30  $\mu\text{mol}$ ) DIC in 150  $\mu\text{l}$  DMF were added and the reaction mixture was stirred at room temperature for 20 min, the product purified by RP-HPLC and lyophilized.

The lyophilized product was incubated for 10 min in 95/5 (v/v) TFA/water and then the solvent was removed in a stream of nitrogen. Product **34** was purified by RP-HPLC and lyophilized (overall yield for all three steps: 20 mg (47%)).

15 HPLC and lyophilized (overall yield for all three steps: 20 mg (47%)).

MS: 17700 – 18200 (broad peak) (MW calculated = 17749 g/mol)

#### Synthesis of compound **9h**

20

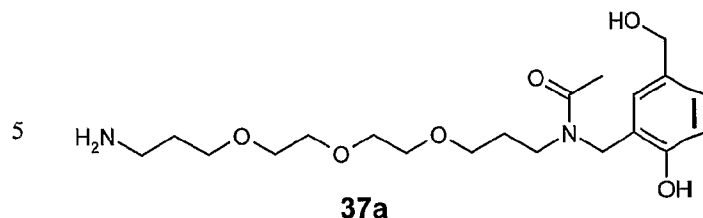
1.5 mg (225 nmol) **8g** and 5 mg (280 nmol) **34** were mixed, dissolved in 300  $\mu\text{l}$  2/1 (v/v) 0.1 M sodium phosphate buffer pH 7/acetonitrile and incubated for 15 min at room temperature. The product was purified by RP-HPLC and lyophilized. (yield 4 mg, 160 nmol, 70%)

25

The lyophilized product was dissolved in 200  $\mu\text{l}$  0.1 M sodium citrate buffer pH 1.5 and 69 mg (20.5  $\mu\text{mol}$ ) **33** in 200  $\mu\text{l}$  2/1 (v/v) acetonitrile/sodium citrate buffer pH 1.5 were added. The mixture was stirred at room temperature for 24 h and product **9h** was purified by size exclusion chromatography (column: Superdex 200, buffer: 10 mM HEPES pH 7.4, 0.005% Tween-20, 3mM EDTA, flow rate: 0.75 ml/min)

30

SEC elution time: 15 min

Synthesis of compound 37a

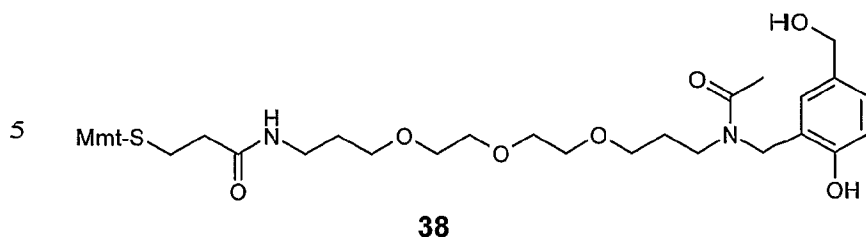
250 mg (0.35 mmol) 2-chlorotrityl chloride resin (loading 1.4 mmol/g) was incubated  
 10 for 1.5 h with 308 mg (4 eq., 1.4 mmol) 4,7,10-trioxatridecane-1,13-diamine in 4 ml  
 DCM to yield **35a**. The resin was washed with DCM and dried. 107 mg (0.7 mmol)  
 HOBt, 110  $\mu$ l (0.7 mmol) DIC, and 150 mg (0.9 mmol) 5-formyl salicylic acid in 3 ml  
 DMF were added and the resulting suspension was stirred for 1 h at RT to yield **36a**.  
 After washing with DCM and THF, the resin was suspended in 6 ml THF and 3 ml (3  
 15 mmol)  $\text{BH}_3$  THF (1 M in THF, 8.5 eq.) were added dropwise. The reaction mixture  
 was stirred for 18 h at 45°C under nitrogen atmosphere. After cooling 4 ml THF, 0.8  
 ml DIEA and 1.6 ml MeOH were added successively. 210 mg (0.84 mmol)  $\text{I}_2$  (as a  
 concentrated THF solution) were added and the suspension was stirred for 1 h. The  
 resin was repeatedly washed (three times each) with THF, DMF, MeOH, and DCM.  
 20 The dried resin was reacted with 107 mg (0.7 mmol) HOBt, 110  $\mu$ l (0.7 mmol) DIC,  
 and 55  $\mu$ L (0.9 mmol) AcOH in 3 ml DMF for 1h. After washing of the resin with  
 DMF and DCM compound **37a** was cleaved from resin with 2/1 (v/v) HFIP/DCM  
 (two times for 30 min). The volatile components were evaporated and the product **37a**  
 was used in the following step without further purification.

25

**37a:** Yield 29 mg (20%) as TFA salt

MS  $[\text{M}+\text{Na}]^+ = 421.4$  (MW+Na calculated = 421.5 g/mol)

30

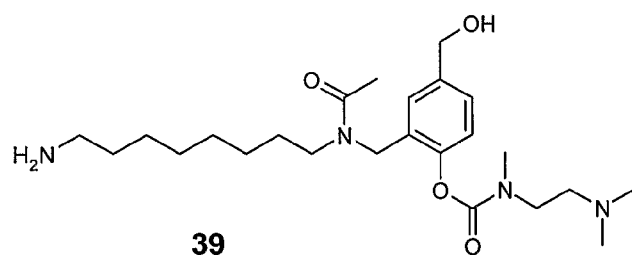
Synthesis of compound 38

24 mg (0.06 mmol) **37a**, 31 mg (0.06 mmol) PyBOP, 32  $\mu$ l (0.18 mmol) DIEA, and  
 10 23 mg (0.06 mmol) **2** in 0.5 ml DMF were reacted for 50 min at room temperature.  
 After addition of 50  $\mu$ l acetic acid product **38** was purified by RP-HPLC.

**38:** Yield 7 mg (15%)

MS  $[M+Na]^+ = 781.3$  (MW+Na calculated = 781.6 g/mol)

15

Synthesis of compound 39

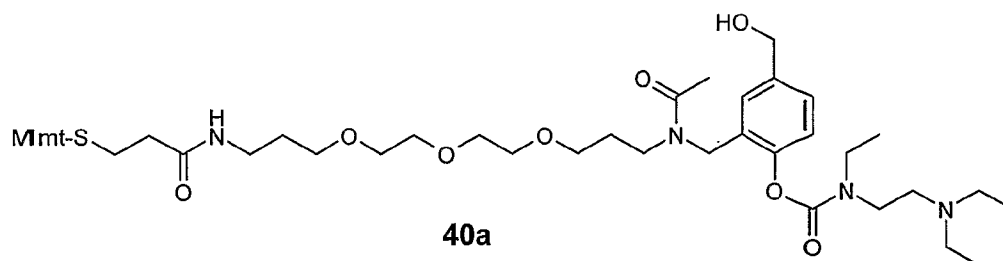
20 300 mg (0.42 mmol) 2-chlorotriyl chloride resin (loading 1.4 mmol/g) was incubated  
 for 1.5 h with 245 mg (4 eq., 1.7 mmol) 1,8-diaminooctane in 4 ml DCM to yield **35b**.  
 The resin was washed with DCM and dried. 107 mg (0.7 mmol) HOBt, 110  $\mu$ l (0.7  
 mmol) DIC, and 150 mg (0.9 mmol) 5-formyl salicylic acid in 3 ml DMF were added  
 and the resulting suspension was stirred for 1 h at RT to yield **36b**. After washing with  
 25 DCM and THF, the resin was suspended in 6 ml THF and 3 ml (3 mmol)  $BH_3$  THF (1  
 M in THF) were added dropwise. The reaction mixture was stirred for 18 h at 45°C  
 under nitrogen atmosphere. After cooling 4 ml THF, 0.8 ml DIEA and 1.6 ml MeOH  
 were added successively. 210 mg (0.84 mmol)  $I_2$  (as a concentrated THF solution)  
 were added and the suspension was stirred for 1 h. The resin was repeatedly washed

(three times each) with THF, DMF, MeOH, and DCM. The dried resin was reacted with 107 mg (0.7 mmol) HOBT, 110  $\mu$ l (0.7 mmol) DIC, and 55  $\mu$ L (0.9 mmol) AcOH in 3 ml DMF for 1h. After washing of the resin with DMF and DCM compound **37b**, 78 mg (0.39 mmol) p-nitrophenylchloroformate, and 210  $\mu$ l (1.2 mmol) DIEA in 1/1 (v/v) THF/DCM were reacted for 30 min at RT. The separated resin was suspended in 1/1 (v/v) THF/DCM and 210  $\mu$ l (1.2 mmol) N,N,N'-trimethylethylenediamine were added. The resulting suspension was stirred for 25 min at RT. The resin was separated and washed with DCM. Product **39** was cleaved from resin with 2/1 (v/v) HFIP/DCM (two times for 30 min). The volatile components were evaporated and the product **39** was purified by HPLC.

**39:** Yield 16 mg (8%) as TFA salt

MS  $[M+Na]^+ = 473.5$  (MW+Na calculated = 473.3 g/mol)

#### Synthesis of compound 40a



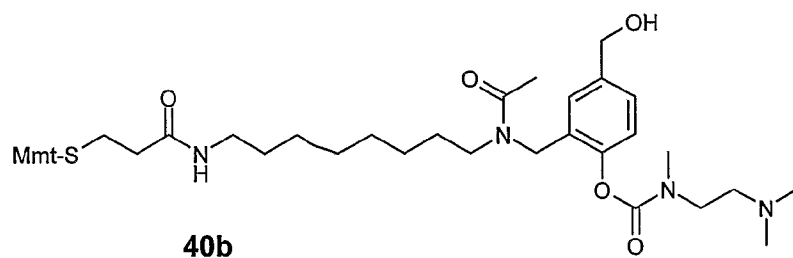
**38** (7 mg, 9  $\mu$ mol) was dissolved in 200  $\mu$ l of dry THF. p-Nitrophenylchloroformate (2.0 mg, 10  $\mu$ mol) and DIEA (4.4  $\mu$ l, 25  $\mu$ mol) were added and the mixture was stirred for 30 min at RT. N,N,N'-Triethylethylenediamine (**15b**) (18  $\mu$ l, 0.1 mmol) was added and stirring was continued for 30 min. The solvent was removed *in vacuo*, 10  $\mu$ l of AcOH were added and **40a** was purified by RP-HPLC.

**40a:** Yield 1 mg (11%) as TFA salt

MS  $[M+Na]^+ = 951.1$  (MW+Na calculated = 951.8 g/mol)



### Synthesis of compound 40b

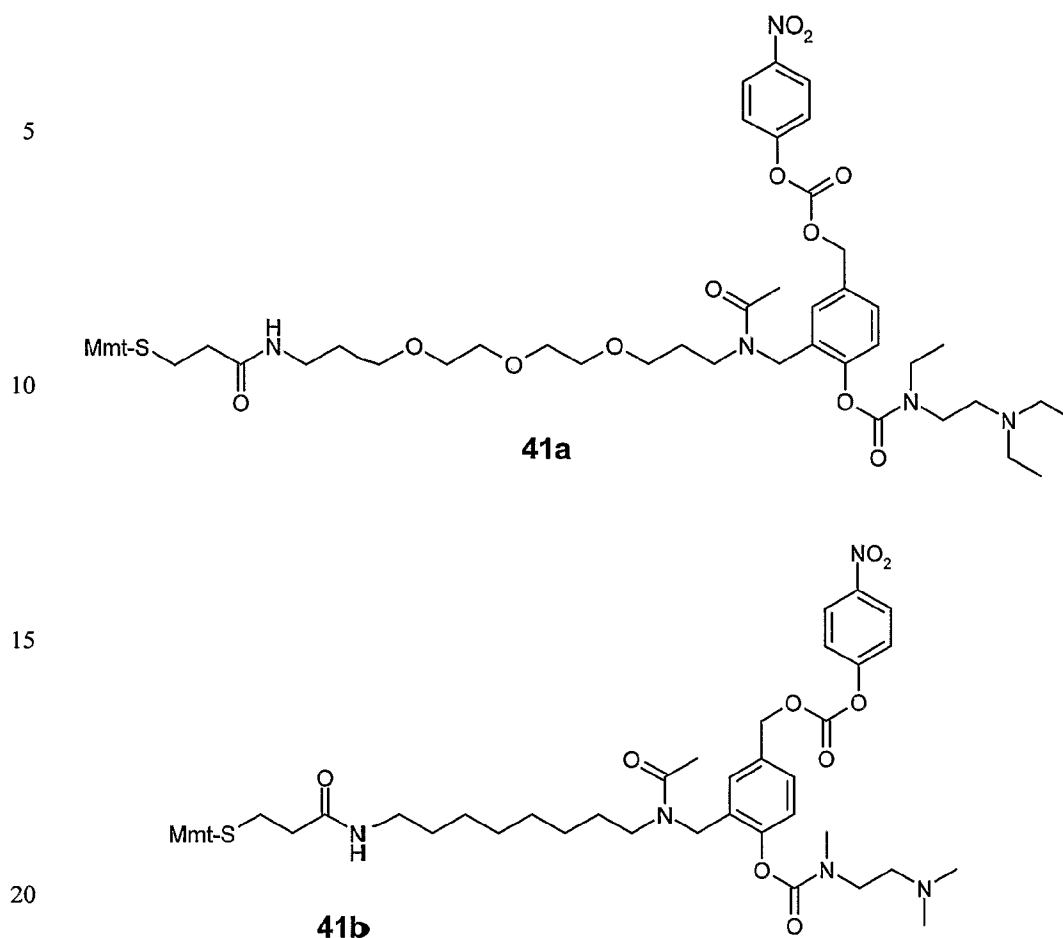


5

15 mg (33  $\mu$ mol) **39**, 18 mg (33  $\mu$ mol) PyBOP, 23  $\mu$ l (0.13 mmol) DIEA, and 13 mg (35  $\mu$ mol) **2** in 0.5 ml DMF were reacted for 45 min at room temperature. After addition of 50  $\mu$ l acetic acid product **40** was purified by RP-HPLC.

10     **40b:** Yield 10 mg (37%) as TFA salt

MS  $[M+H]^+ = 811.5$  (MW+Na calculated = 810.5 g/mol)

Synthesis of compound **41a** and **41b**

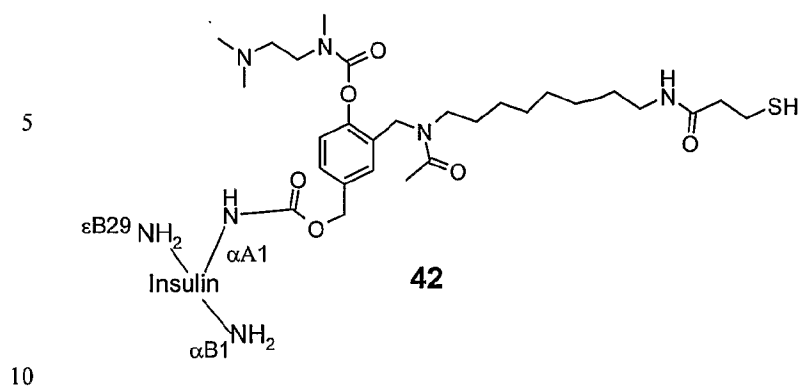
Carbonate **41a** or **41b** was synthesized from **40a** or **40b** as described for compound **7a**

**41a**: Yield 0.4 mg as TFA salt

MS  $[M+Na]^+ = 1116.8$  (MW+Na calculated = 1116.9 g/mol)

**41b**: Yield 2 mg (16%) as TFA salt

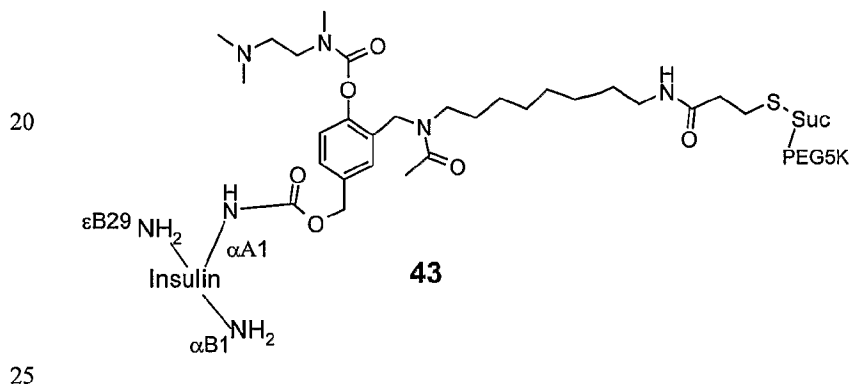
MS  $[M+H]^+ = 976.8$  (MW calculated = 975.8 g/mol)

Synthesis of compound 42

Insulin derivative **42** was synthesized from **41b** as described for compound **8a**.

**42** MS  $[M+3H]^{3+} = 2124.5$   $[M+4H]^{4+} = 1594.6$  (MW calculated = 6371 g/mol)

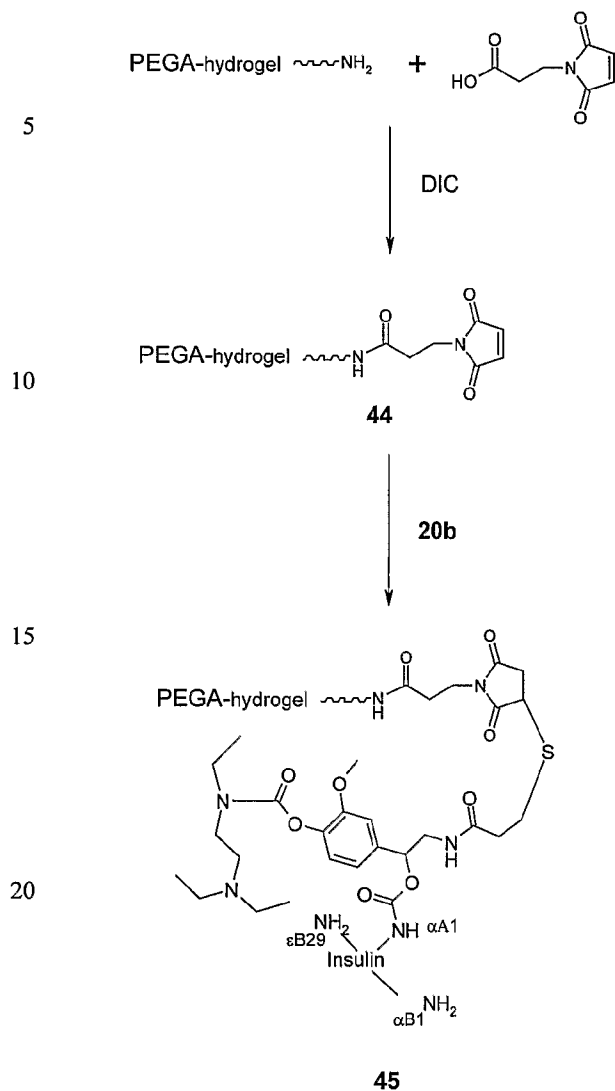
15

Synthesis of compound 43

Insulin derivative **43** was synthesized from **42** as described for compound **9a**.

**43**: SEC retention time: 18.0 min

30

Synthesis of rh-insulin loaded PEGA hydrogel 45Maleimide derivatization of polyacrylamide based hydrogel (PEGA):

NH<sub>2</sub>-PEGA hydrogel beads with 0.4 mmol/g loading and 150-300  $\mu\text{m}$  bead size were purchased from Novabiochem.

- 2.5 g methanol-wet NH<sub>2</sub>-PEGA-hydrogel (0.4 mmol/g NH<sub>2</sub>-loading) was weighed into a syringe equipped with a polypropylene frit. Maleimide loading was adjusted by acylation employing a mixture of activated maleimidopropionic acid and acetic acid as described in the following. The hydrogel was washed 5 times with DMF and reacted with 13.5 mg (0.08 mmol) 3-maleimidopropionic acid, 115.2  $\mu\text{l}$  (1.92 mmol)

acetic acid and 313  $\mu$ l (2 mmol) DIC in 4 ml DMF for 30 min. The maleimide derivatized hydrogel **44** was washed 10 times with DMF and DCM and finally with acetonitrile.

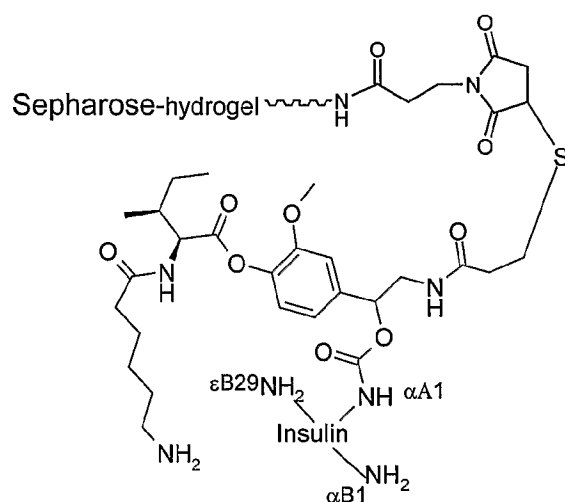
- 5 30 mg of maleimide derivatized resin **44** (loading 16  $\mu$ mol/g) was reacted with 3 mg of compound **20b** (480 nmol, 1.06 eq) in 600  $\mu$ l 20/80 (v/v) acetonitrile/50 mM phosphate buffer (pH 7.4) for 10 min to give rh-insulin loaded hydrogel **45**. The hydrogel **45** was washed 5 times with 50/50 (v/v) acetonitrile/water and three times with acetonitrile and dried under vacuum.

10

#### Synthesis of rh-insulin carbohydrate-based hydrogel **46**

15

20

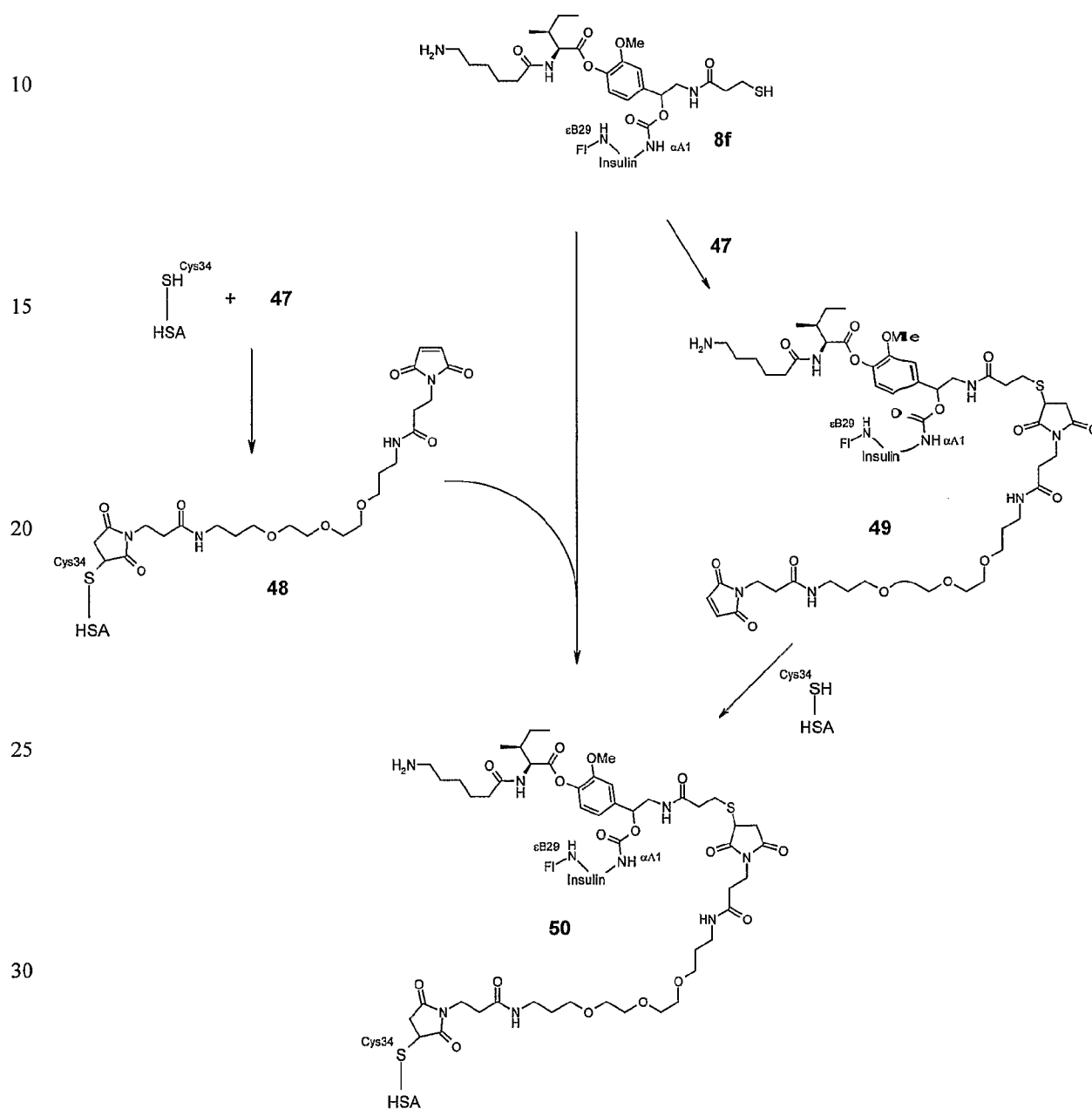
**46**

- 25 NHS-activated "Sepharose 4 Fast Flow" hydrogel beads (chemically crosslinked agarose, crosslinker epichlorhydrin) were purchased from Amersham. 1.5 g ethanol-wet Sepharose hydrogel (150 mg dry hydrogel) was weighed into a syringe equipped with a polypropylene frit and reacted with 1 M 4,7,10-trioxatridecane-1,13-diamine in DMF for 30 min. After 5 washing steps with DMF,
- 30 hydrogel was reacted with 8.5 mg (0.05 mmol) 3-maleimidopropionic acid, 57  $\mu$ l (0.95 mmol) acetic acid, 151 mg (1 mmol) HOBt and 158  $\mu$ l (1 mmol) DIC in 4 ml DMF for 30 min to give maleimide derivatized hydrogel. The hydrogel was washed 10 times with DMF and finally with acetonitrile.

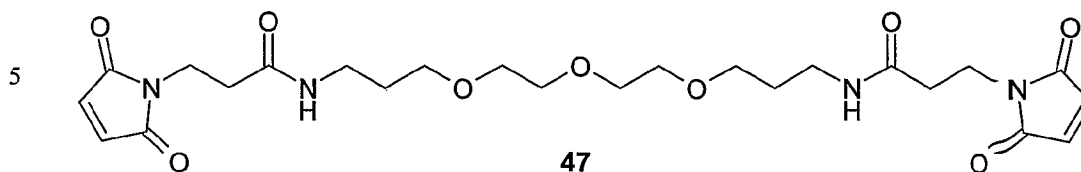
1.5 mg **8c** was dissolved in 25/75 (v/v) acetonitrile/50 mM phosphate buffer pH 7.4 and reacted with 10.8 mg maleimide derivatized hydrogel for 10 min. The rh-insulin loaded hydrogel **46** was washed five times with 50/50 (v/v) acetonitrile/water and three times with acetonitrile and dried under vacuum.

5

### Synthesis scheme of Fluorescein-Insulin-rHSA (50)



### Synthesis of bismaleimide **47**



3-Maleimido-propionic acid (92 mg, 0.54 mmol) in 200  $\mu$ l DMF was reacted with  
DIC (78  $\mu$ l, 0.50 mmol) at RT for 15 min. 4,7,10-Trioxa-tridecan-1,13-diamine (43.5  
10  $\mu$ l, 0.20 mmol) was added and the mixture was stirred for 30 min at RT.  
After addition of 800  $\mu$ l 1/4 (v/v) acetic acid/water **47** was purified by RP-HPLC.

47: Yield 23 mg (22%)

MS  $[M+Na]^+ = 545.5$  (MW+Na calculated = 545.6 g/mol)

### Synthesis of rHSA-maleimide (48)

66.5  $\mu$ l 3 mM rHSA solution in 145 mM NaCl, 32 mM sodium octanoate, 0.015% Tween-80 was mixed with 66.5  $\mu$ l 0.5 M phosphate buffer pH 7.0. 0.41 mg bismaleimide **47** (0.8  $\mu$ mol) were added and the mixture was reacted for 15 min at RT. Compound **48** was purified by SEC (column: Superdex 200, flow rate: 0.75 ml/min) using 10 mM HEPES buffer pH 7.4, 150 mM NaCl, 3 mM EDTA, and 0.005% Tween as mobile phase. (Yield: 2.6 ml 77.5  $\mu$ M **48**)

25 SEC retention time: 17.1 min (280 nm)

ESI-MS = 66988 (MW calculated = 66984 g/mol)

### Synthesis of fluorescein-insulin-linker-maleimide (49)

30 40  $\mu$ l of 2.4 mM bismaleimide **47** in 1/1 (v/v) acetonitrile/water (96 nmol) were mixed with 40  $\mu$ l 0.5 M sodium borate buffer pH 5.8. 24 nmol **8f** in 16.8  $\mu$ l 1/1 (v/v) acetonitrile/water were added and the mixture was incubated for 10 min at RT. 5  $\mu$ l AcOH were added and **49** was purified by RP-HPLC.

ESI-MS = 7211 (MW calculated = 7211 g/mol)

Synthesis of fluorescein-insulin-linker-rHSA **50**

a) from **49** and rHSA

5 b) from **48** and **8f**

a)

30  $\mu$ l of 80  $\mu$ M **49** in 1/1 (v/v) acetonitrile/water (2.4 nmol) were mixed with 70  $\mu$ l  
0.25 M sodium phosphate buffer pH 6.4. 8  $\mu$ l 3 mM rHSA in 145 mM NaCl 32 mM  
10 sodium octanoate, 0.0015% Tween-80, (24 nmol) was added and the mixture was  
incubated at RT for 20 min.

Compound **50** was purified by SEC (column: Superdex 200, flow rate: 0.75 ml/min)  
using 10 mM HEPES buffer pH 7.4, 150 mM NaCl, 3 mM EDTA, and 0.005%  
Tween as mobile phase.

15

SEC retention time: 17.3 min (500 nm)

ESI-MS = 73676 (MW calculated = 73673 g/mol)

b)

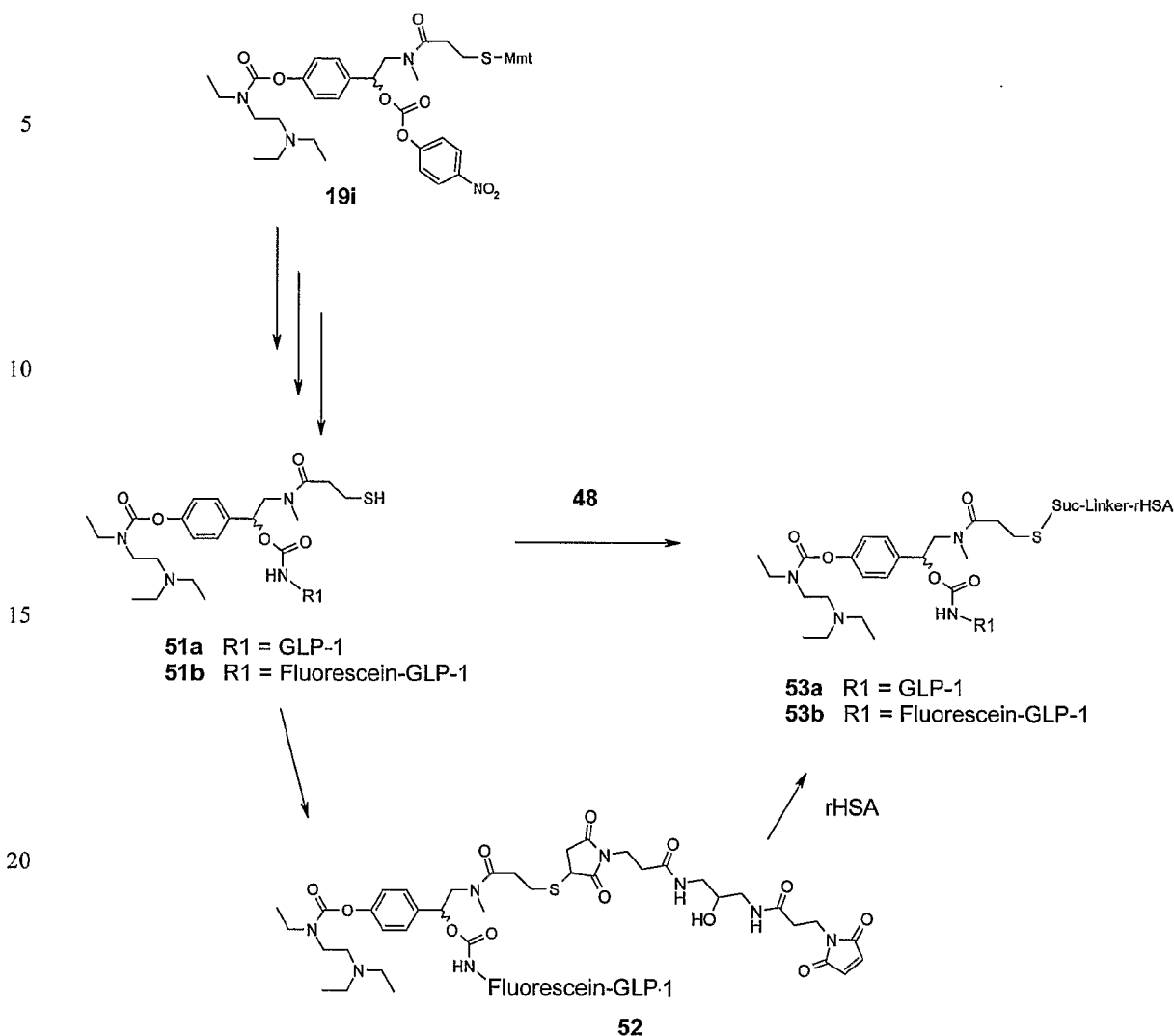
20 SEC eluate of rHSA-maleimide **48** (241  $\mu$ l, 77.5  $\mu$ M, 18.7 nmol) was mixed with 20  
 $\mu$ l 0.5 M sodium borate buffer pH 5.8. 14  $\mu$ l 1.41 mM **8f** (19.6 nmol) in 1/1 (v/v)  
acetonitrile/water were added and the mixture was incubated at RT for 10 min. 1.2  $\mu$ l  
48.5 mM 3-maleimido propionic acid (58 nmol) in 1/1 (v/v) acetonitrile/water were  
added and compound **50** was purified by SEC (column: Superdex 200, flow rate: 0.75  
25 ml/min) using 10 mM HEPES buffer pH 7.4, 150 mM NaCl, 3 mM EDTA, and  
0.005% Tween as mobile phase.

SEC retention time: 17.1 min (500 nm)

ESI-MS = 73698 (MW calculated = 73673 g/mol)

30



Synthesis scheme of rHSA-linker-GLP-1 **53a** and **53b**Synthesis of **51a**

GLP(7-36) (sequence: HAEGTFTSDVSSYLEGQAAKEFIAWLVKGR-amide) was synthesized on Rink-amide resin employing fmoc-strategy (Specialty Peptide

30 Laboratories, Heidelberg, Germany). N-terminal fmoc-protecting group was removed and the resin was washed with DCM and dried. 118 mg resin (0.11 mmol/g, 13.2  $\mu$ mol) was suspended in a solution of 50 mg **19i** (53  $\mu$ mol) in 750  $\mu$ l dry DMSO and 22.4  $\mu$ l DIEA. 2.1  $\mu$ l pyridine was added and the mixture was shaken for 48 h at RT. After washing the resin 6 times each with DMF and DCM, cleavage of the peptide

from resin and removal of protecting groups was achieved with 96/2/2 (v/v/v) TFA/triethylsilane/water. Volatiles were removed under nitrogen flow and **51a** was purified by RP-HPLC and lyophilized.

5 **51a:** Yield 4.6 mg (9%)

MS:  $[M+3H]^{3+} = 1251.0$  (MW calculated = 3750.3 g/mol)

#### Synthesis of **51b**

Lys28 ivDde side chain protected GLP(7-36) (sequence:

- 10 HAEGTFTSDVSSYLEGQAAKEFIAWLVK(ivDde)GR-amide) was synthesized on Rink-amide resin employing fmoc-strategy (Specialty Peptide Laboratories, Heidelberg, Germany). N-terminal fmoc-protecting group was removed and the resin was washed with DCM and dried. 50 mg resin (0.11 mmol/g, 5.5  $\mu$ mol) was suspended in a solution of 25 mg **19i** (26  $\mu$ mol) in 400  $\mu$ l dry DMSO and 11.2  $\mu$ l
- 15 DIEA. 1.1  $\mu$ l pyridine was added and the mixture was shaken for 48 h at RT. After washing the resin six times with DMF the ivDde protecting group was cleaved by incubating the resin 3 times with 5% hydrazine in DMF for 20 min. Fmoc-8-amino, 3,6-dioxaoctanoic acid was coupled according to the standard coupling cycle. Fmoc protecting group was removed and carboxy-fluorescein was coupled by incubating the
- 20 resin with 8 mg 5-(and-6)-carboxyfluorescein succinimidyl ester and 2  $\mu$ l DIEA for 60 min. Resin was washed six times each with DMF and DCM. Cleavage of the peptide from resin and removal of protecting groups was achieved with 96/2/2 (v/v/v) TFA/triethylsilane/water. Volatiles were removed under nitrogen flow. **51b** was used for the synthesis of **52** without further purification.

25 MS:  $[M+3H]^{4+} = 1064.3$ ,  $[M+2H]^{3+} = 1418.3$  (MW calculated = 4254 g/mol)

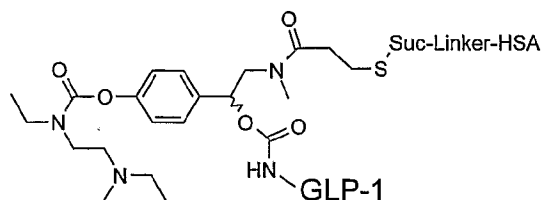
#### Synthesis of **52**

Raw material **51b** was dissolved in 500  $\mu$ l 1/1 (v/v) acetonitrile / 0.25 M sodium phosphate pH 7 and 8 mg N,N'-bis(3-maleimidopropionyl)-2-hydroxy-1,3-

- 30 propanediamine were added. The solution was stirred at RT for 15 min and **52** was purified by RP-HPLC and lyophilized.

**52:** Yield: 5.1 mg

MS  $[M+3H]^{4+} = 1162.8$ ,  $[M+2H]^{3+} = 1549.4$  (MW calculated = 4645 g/mol)

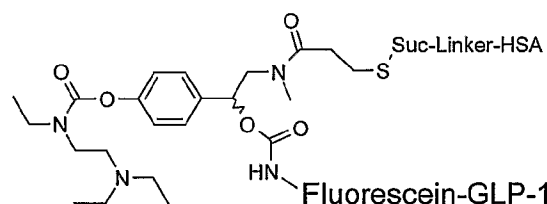
Synthesis of compound 53a**53a**

- 5 30  $\mu$ l 1.57 mM **48** (47 nmol) in 10 mM HEPES buffer pH 7.4, 150 mM NaCl, 3 mM EDTA, and 0.005% Tween were mixed with 10  $\mu$ l 0.5 M sodium phosphate buffer pH 7.4. A mixture of 2  $\mu$ l DMSO and 12  $\mu$ l of 6.06 mM **51a** (73 nmol) in water/acetonitrile 9/1 (v/v) was added and the solution was incubated at RT for 30 min. **53a** was purified by SEC (column: Superdex 200, flow rate: 0.75 ml/min) using
- 10 10 mM phosphate buffer pH 7.4, 150 mM NaCl, and 0.005% Tween as mobile phase.

SEC retention time: 17.7 min (280 nm)

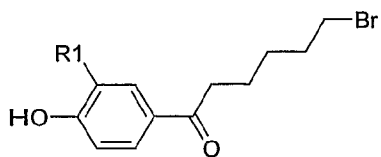
ESI-MS = 70745 (MW calculated = 70734 g/mol)

- 15 Synthesis of **53b**

**53b**

- 100  $\mu$ l 3 mM **52** (300 nmol) in 9/1 50 mM sodium phosphate pH 7.0 / acetonitrile were mixed with 100  $\mu$ l 3 mM HSA (300 nmol) and the solution was incubated at RT
- 20 for 30 min. **53b** was purified by SEC (column: Superdex 200, flow rate: 0.75 ml/min) using 10 mM HEPES buffer pH 7.4, 150 mM NaCl, 3 mM EDTA, and 0.005% Tween as mobile phase.

SEC retention time: 17.7 min (500 nm)

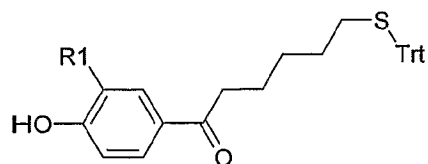
Synthesis of compounds **54a** and **54b****54a** R1 = Me5 **54b** R1 = H

AlCl<sub>3</sub> (1.05 eq) was suspended in DCM and 6-bromohexanoic acid chloride (1 eq) was added. After stirring at RT for 20 min o-cresol (1 eq) was added and the mixture was reacted at RT for 25 min. The reaction mixture was poured into ice water and  
 10 extracted with ethyl acetate. The separated organic phase was dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated *in vacuo*. Product **54a** was purified by silica gel column chromatography using heptane/ethyl acetate (4/1) as mobile phase.

**54b** was synthesized as described above using 6-bromohexanoic acid chloride and  
 15 phenol.

**54a:** Yield 3.7 g (33%)MS [M+H]<sup>+</sup> = 285.1 and 287.2 (MW+H calculated = 386.2 g/mol)**54b:** Yield 620 mg (15%)MS [M+H]<sup>+</sup> = 271.2 (MW calculated = 271.0 g/mol)

20

Synthesis of compounds **55a** and **55b**25 **55a** R1 = Me**55b** R1 = H

DBU (105  $\mu$ l, 701  $\mu$ mol) was added to a solution of bromide **54a** (105 mg, 369  $\mu$ mol) and tritylthiol (204 mg, 738  $\mu$ mol) in 50 ml dry DMSO. The reaction mixture was stirred at RT for 40 min and acidified with 1 N H<sub>2</sub>SO<sub>4</sub>. The aqueous layer was extracted with ethyl acetate and evaporated. **55a** was purified by RP-HPLC.

- 5 **55b** was synthesized according to the same protocol using **54b** (180 mg, 0.66 mmol).

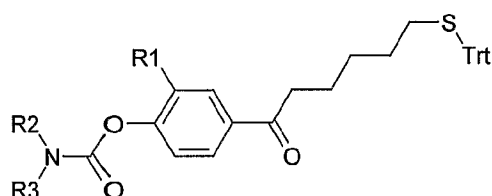
**55a**: Yield 173 mg (97%)

MS [M+Na]<sup>+</sup> = 503.6 (MW+Na calculated = 503.7 g/mol)

**55b**: Yield 160 mg (85%)

- 10 MS [M+Na]<sup>+</sup> = 489.5 (MW+Na calculated = 489.3 g/mol)

#### Synthesis of compounds **56a** and **56b**



- 15 **56a** R1 = Me, R2 = R3 = 3-(dimethylamino)propyl,  
**56b** R1 = H, R2 = Me, R3 = 3-(N-ethyl-N-methylamino)propyl

**56a** was prepared from **55a** (9 mg, 19  $\mu$ mol), p-nitrophenyl-chloroformate and bis(3-dimethylamino-propyl)amine (21  $\mu$ l, 94  $\mu$ mol) as described for compound **16a**.

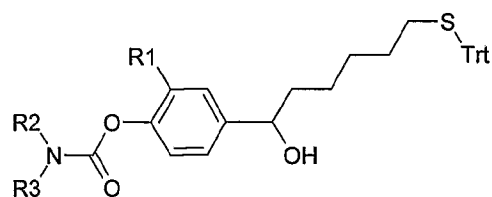
- 20 **56b** was synthesized from **55b** (160 mg, 0.34 mmol), p-nitrophenyl-chloroformate and N-ethyl-N,N'-dimethyl-1,3-propanediamine (**15c**) as described for compound **16a**.

**56a**: Yield 12 mg (70%) as TFA salt

MS [M+Na]<sup>+</sup> = 716.8 (MW+Na calculated = 717.0 g/mol)

- 25 **56b**: Yield 80 mg (32%) as TFA salt

MS [M+Na]<sup>+</sup> = 645.6 (MW+Na calculated = 645.4 g/mol)

Synthesis of compounds 57a and 57b

5

**57a** R1 = Me, R2 = R3 = 3-dimethylamino-propyl,

**57b** R1 = H, R2 = Me, R3 = 3-(N-ethyl-N-methylamino)propyl

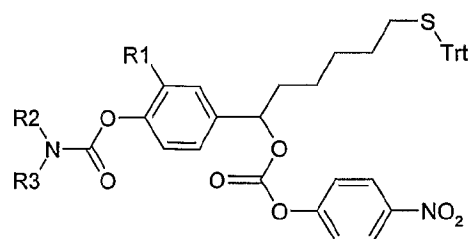
**57a** and **57b** were synthesized from **56a** (12 mg, 13  $\mu$ mol, double TFA salt) and **56b**  
 10 (80 mg, 110  $\mu$ mol, TFA salt), respectively, as described for compound **16g**.

**57a:** Yield 9 mg (75%) as TFA salt

MS  $[M+Na]^+ = 719.0$  (MW+Na calculated = 718.7 g/mol)

**57b:** Yield 60 mg (75%) as TFA salt

15 MS  $[M+Na]^+ = 647.4$  (MW+Na calculated = 647.4 g/mol)

Synthesis of compounds 58a and 58b

20

**58a** R1 = Me, R2 = R3 = 3-(dimethylamino)propyl,

**58b** R1 = H, R2 = Me, R3 = 3-(N-ethyl-N-methylamino)propyl

25 **57a** (1 eq, 8mg, 9  $\mu$ mol), 4-nitrophenyl chloroformate (3.5 eq, 6 mg, 30  $\mu$ mol), DIEA  
 (6 eq, 9  $\mu$ l, 52  $\mu$ mol), and DMAP (1 eq, 1 mg, 9  $\mu$ mol) were stirred in 1 ml dry DCM

at RT for 45 min under nitrogen atmosphere. The volatiles were evaporated and acetic acid was added. The mixture was dissolved in 1/1 (v/v) acetonitrile/water and the carbonate **58a** was purified by RP-HPLC.

Carbonate **58b** was prepared likewise from **57b** (135 mg, 0.18 mmol).

5

**58a**: Yield 7 mg (70%) as TFA salt

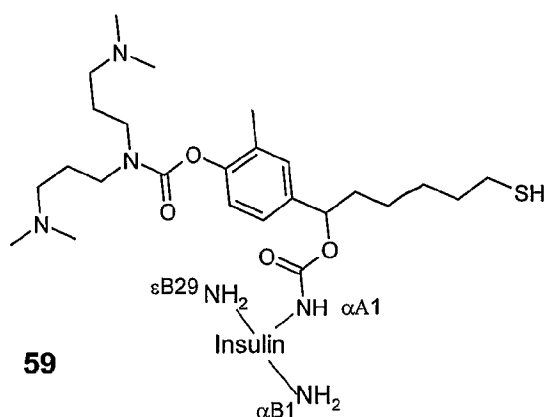
MS  $[M+Na]^+ = 883.8$  (MW+Na calculated = 884.1 g/mol)

**58b**: Yield 110 mg (77%) as TFA salt

MS  $[M+Na]^+ = 812.4$  (MW+Na calculated = 812.5 g/mol)

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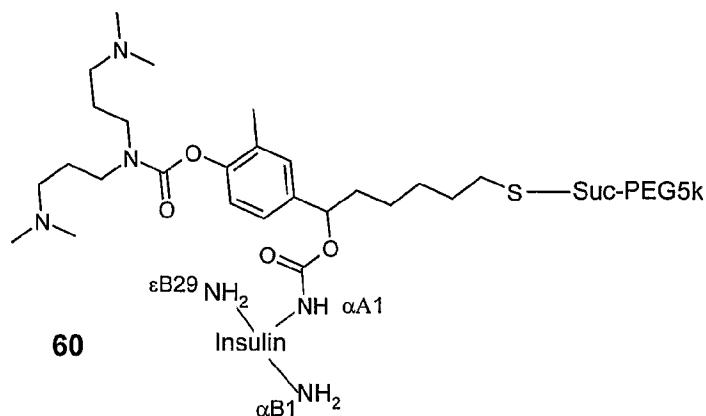
### Synthesis of compound **59**



15 Rh-Insulin (44.5 mg, 7.7  $\mu$ mol), carbonate **58a** (1 eq, 7 mg, 6.4 mmol), DIEA (15  $\mu$ l, 88  $\mu$ mol) and DMAP (1.5 mg, 12  $\mu$ mol) in 0.3 ml DMSO were reacted at RT for 30 min. The reaction mixture was neutralized with acetic acid and diluted with 1/1 (v/v) acetonitrile/water. RP-HPLC purification gave the appropriate Trt-protected intermediate.

20 After lyophilization, the Trt-protected intermediate was mixed with 95/5 (v/v) TFA/triethylsilane and stirred for 5 min. Volatiles were removed under nitrogen flow and **59** was purified by RP-HPLC and lyophilized. Position of insulin modification was verified by DTT reduction and MS analysis.

25 **59**: MS  $[M+3H]^{3+} = 2095.5$   $[M+4H]^{4+} = 1572.2$  (MW calculated = 6288 g/mol)

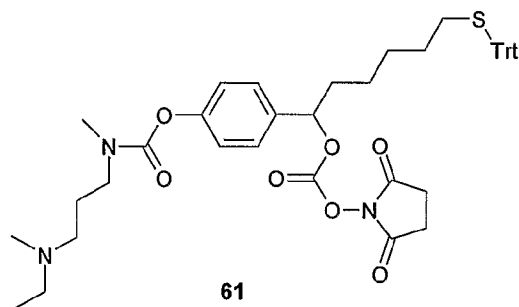
Synthesis of compound 60

5 **60** was prepared from **59** (0.17  $\mu\text{mol}$ ) as described for compound **9a**.

**60**: SEC retention time: 19.5 min

Synthesis of compound 61

10



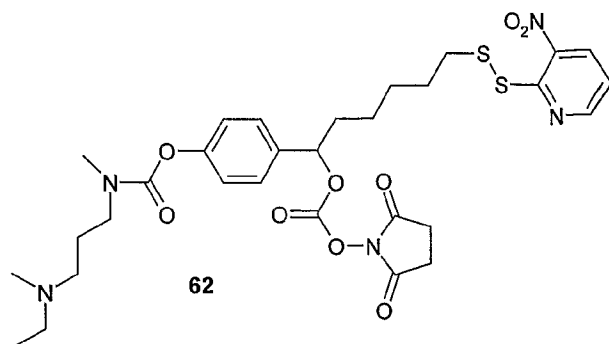
**57b** (70 mg, 90  $\mu\text{mol}$ ), DSC (161 mg, 630  $\mu\text{mol}$ ), DIEA (192  $\mu\text{l}$ , 1.1 mmol), and DMAP (11 mg, 90  $\mu\text{mol}$ ) were stirred in 1 ml dry acetonitril at RT for 14 h under nitrogen atmosphere. The volatiles were evaporated and acetic acid was added. The mixture was dissolved in 1/1 (v/v) acetonitrile/water and the carbonate **61** was purified by RP-HPLC.

**61**: Yield 40 mg (51%) as TFA salt

20 MS  $[\text{M}+\text{Na}]^+ = 788.4$  (MW+Na calculated = 788.5 g/mol)



### Synthesis of compound 62

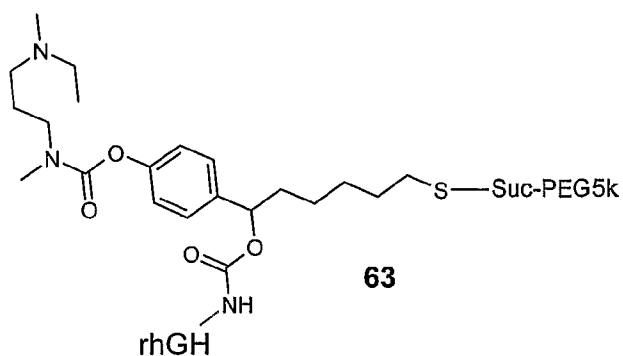


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**61** (12 mg, 13  $\mu\text{mol}$ ) and NPys-Cl (4 mg, 21  $\mu\text{mol}$ ) were stirred in 1 ml DCM at -10°C for 2 h under nitrogen atmosphere. Volatiles were removed under nitrogen flow and **62** was purified by RP-HPLC.

**62:** Yield 7 mg (65%) as TFA salt

MS  $[M+Na]^+ = 700.9$  (MW+Na calculated = 701.4 g/mol)

15 Synthesis of compound 63

0.9 mg desalted rhGH (ProspecTany, Israel, MW 22250 g/mol, 40 nmol) in 200  $\mu$ l 50  
20 mM borate buffer (pH 8.0), 8  $\mu$ l of carbonate **62** in acetonitrile (38 mM, 300 nmol),  
and 40  $\mu$ l DMSO were reacted at RT for 3 h. The solvent mixture and low molecular

weighed compounds were replaced by water and subsequently by acetate buffer (25 mM, pH 4.2, 0.005% Tween 20) by ultrafiltration using Centricon 5 filter (cutoff 5 kDa). 8  $\mu$ l (80 nmol) 10 mM DTT in 25 mM acetate buffer pH 4.2, 0.005% Tween was added and incubated at RT for 30 min. Low molecular weight compounds were removed by ultrafiltration using Centricon 5 filter and 25 mM acetate buffer pH 4.2, 0.005% Tween as eluate. After concentration to a volume of 100  $\mu$ l (Centricon 5) 20  $\mu$ l (100 nmol) 5 mM maleimide-PEG5k in water and 80  $\mu$ l 0.5 M phosphate buffer pH 7.0 were added. The mixture was incubated at RT for 5 min. Monoconjugate **63** was separated by SEC (column: Superdex 200, flow rate: 0.75 ml/min) using 10 mM phosphate buffer pH 7.4, 150 mM NaCl, and 0.005% Tween 20 as mobile phase. The collected eluate (approximately 1.0 ml) was diluted with 0.5 ml buffer containing 0.05% NaN<sub>3</sub> and directly used for release rate determination.

**63**: SEC retention time: 17.5 min

#### Release of insulin or fluorescein-insulin from conjugates in buffer pH 7.4

Release of (fluorescein)-insulin from (fluorescein)-insulin conjugates **9a** to **9h**, **21a** to **21f**, **30a**, **30b**, **43**, **50**, and **60**, release of fluorescein-GLP-1 from **53b**, and release of rhGH from **63** was effected by linker hydrolysis in aqueous buffer pH 7.4. Collected SEC eluates of (fluorescein)-insulin conjugates (see above), fluorescein-GLP-1 conjugate and rhGH-conjugate, respectively, were incubated at 37 °C and samples were taken at time intervals and analyzed by RP-HPLC (insulin conjugates) or SEC (rhGH conjugate, fluorescein insulin conjugates and fluorescein-GLP-1 conjugate) and UV detection at 215 or 280 nm or VIS detection at 500 nm. Peaks correlating with the retention time of native insulin, fluorescein-insulin, fluorescein-GLP-1, and rhGH, respectively, were integrated and plotted against incubation time, and curve-fitting software was applied to estimate the corresponding halftime of release.

#### Release of insulin from hydrogel conjugates **45** and **46**

4 mg of **45** or 2 mg **46** were weighed into a test tube and incubated with 1 ml 10 mM HEPES buffer pH 7.4, 150 mM NaCl, 0.005% Tween at 37°C. 45  $\mu$ l samples were

taken at different time intervals and quantitatively analyzed for rh-insulin by a RP-HPLC assay. The rh-insulin peaks were integrated and rh-insulin concentration was obtained from a standard curve. A first order release kinetic was fitted to the data points to give the linker half life.

5

#### MS-analysis of released insulin from compound 9a, 9b and 30a

Samples of buffer released insulin (see above) were analyzed by mass spectrometry. Figure 11 shows the mass spectra of released insulin from compound 9a, 9b, and 30a.

- 10 The mass spectrum of insulin released from compound 9a clearly shows a major side product (indicated by arrows), corresponding to the irreversibly pentanoyl-modified insulin. In this case, removal of the pentanoyl masking group was not by hydrolysis but by acyl transfer to the insulin. The mass spectrum of insulin released from compound 9b and 30a shows no modification.

15

#### Release of fluorescein-insulin from conjugate 9d and 9e in 80% human plasma

Release of fluorescein-insulin from 9d or 9e was effected by hydrolysis in 80% human plasma in 20 mM HEPES pH 7.4 at 37 °C. Samples were taken at time intervals and analyzed by SEC and VIS detection at 500 nm. Peaks correlating with the retention time of fluorescein-insulin were integrated and plotted against incubation time, and curve-fitting software was applied to estimate the corresponding halftime of release.

- 25 Table: Polymeric prodrug hydrolysis

compound	t <sub>1/2</sub> buffer pH 7.4	t <sub>1/2</sub> human plasma
9a	40 h	nd
9b	55 h	nd
9c	4.5 d	nd
9d	7 h	4 h
9e	55 h	30 h
9f	90 h	nd
9g	37 h	nd

<b>9h</b>	88 h	nd
<b>21a</b>	64 d	nd
<b>21b</b>	8 d	nd
<b>21c</b>	52 d	nd
<b>21d</b>	29 d	nd
<b>21e</b>	100 d	nd
<b>21f</b>	83 h	nd
<b>30a</b>	17 d	nd
<b>30b</b>	> 70 d	nd
<b>43</b>	4 h	nd
<b>45</b>	7 d	nd
<b>46</b>	4 d	nd
<b>50</b>	57 h	nd
<b>53b</b>	19 h	nd
<b>60</b>	10 d	nd
<b>63</b>	51 d	nd

nd = not determined

The foregoing is considered illustrative of the principles of the invention and since numerous modifications will occur to those skilled in the art, it is not intended to limit the invention to the exact construction and operation described. All suitable modifications and equivalents fall within the scope of the claims.

## Abbreviations:

	Boc	t-butyloxy carbonyl
	DBU	1,3-diazabicyclo[5.4.0]undecene
	DCM	dichloromethane
5	(iv)Dde	1-(4,4-dimethyl-2,6-dioxo-cyclohexyliden)3-methyl-butyl
	DIC	diisopropylcarbodiimide
	DIEA	diisopropylethylamine
	DMAP	dimethylamino-pyridine
	DMF	N,N-dimethylformamide
10	DMSO	dimethylsulfoxide
	Dpr	diaminopropionic acid
	DSC	disuccinidylcarbonate
	EDTA	ethylenediaminetetraacetic acid
	Et	ethyl
15	eq	stoichiometric equivalent
	fmoc	9-fluorenylmethoxycarbonyl
	Fmoc-Ado-OH	Fmoc-8-amino-3,6-dioxaoctanoic acid
	HFIP	hexafluoroisopropanol
	HEPES	N-(2-hydroxyethyl) piperazine-N'-(2-ethanesulfonic acid)
20	HOBt	N-hydroxybenzotriazole
	LCMS	mass spectrometry-coupled liquid chromatography
	Mal	maleimidopropionyl
	Me	methyl
	Mmt	4-methoxytrityl
25	MS	mass spectrum
	MW	molecular mass
	Npys	3-nitro-2-pyridinesulfonyl
	PyBOP	benzotriazole-1-yl-oxy-tris-pyrrolidino-phosphonium hexafluorophosphate
30	rHSA	recombinant human serum albumin
	rhGH	recombinant human growth hormone
	RP-HPLC	reversed-phase high performance liquid chromatography
	RT	room temperature

	SEC	size exclusion chromatography
	Suc	succinimidopropionyl
	TES	triethylsilane
	TFA	trifluoroacetic acid
5	THF	tetrahydrofuran
	UV	ultraviolet
	VIS	visual
	Z	benzyloxycarbonyl

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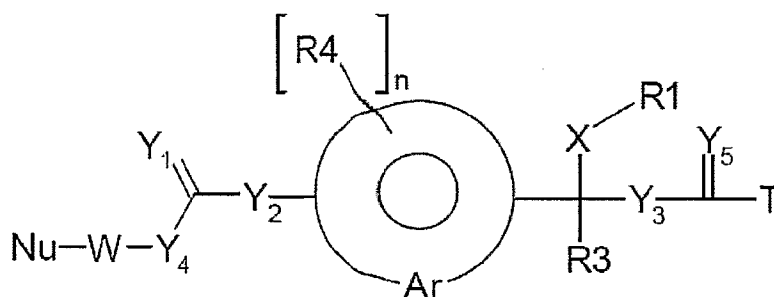
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## CLAIMS

1. A polymeric cascade prodrug comprising:
  - an amine containing biologically active moiety
  - a masking group having at least one nucleophile and being distinct from the carrier.
2. The prodrug of claim 1 or corresponding polymeric cascade prodrug linker reagent having the following structure:



- wherein
- T is D or A
- D being a residue of an amine containing biologically active moiety and
- A being a leaving group;
- X is a spacer moiety such as R5-Y6
- Y<sub>1</sub>, Y<sub>2</sub> can each be either O, S, or NR<sub>6</sub>, independently of each other.
- Y<sub>3</sub>, Y<sub>5</sub> can each be either O or S, independently of each other.
- Y<sub>4</sub> is O, NR<sub>6</sub>, or -C(R<sub>7</sub>)(R<sub>8</sub>)-
- Y<sub>6</sub> is O, S, NR<sub>6</sub>, succinimide, maleimide, unsaturated carbon-carbon bonds or any heteratom containing a free electron pair or is absent..
- R<sub>3</sub> is selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryl, cyano, nitro, halogen, carboxy, carboxyalkyl, alkylcarbonyl, or carboxamidoalkyl;
- R<sub>4</sub> is selected independently from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl, substituted or non-substituted linear, branched, or cyclical alkoxy, substituted or non-substituted linear, branched, or cyclical heteroalkoxy, aryloxy, or heteroaryloxy, cyano, halogen;
- R<sub>5</sub> is selected from substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryl;

R7 and R8 are selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryl, carboxyalkyl, alkylcarbonyl, carboxamidoalkyl, cyano, or halogen;

R6 is selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryl;

R1 is a polymer;

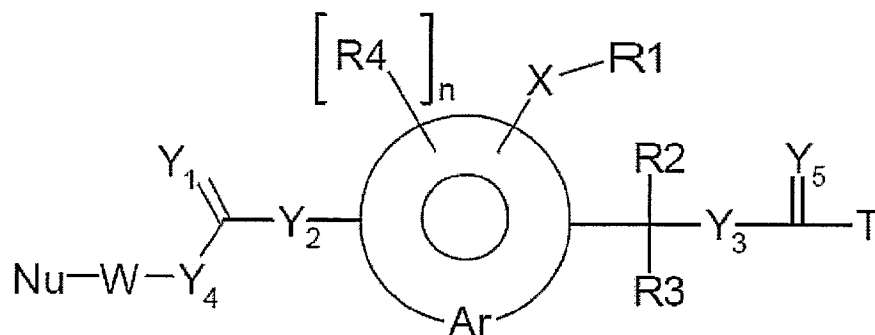
W is selected from substituted or non-substituted linear, branched or cyclical alkyl, aryls, substituted aryls, substituted or non-substituted linear, branched or cyclical heteroalkyl, substituted or nonsubstituted heteroaryl;

Nu is a nucleophile;

n is zero or a positive integer; and

Ar is a multi-substituted aromatic hydrocarbon or a multi-substituted aromatic heterocycle.

3. The prodrug of claim 1 or corresponding polymeric cascade prodrug linker reagent having the following structure:



wherein

T is D or A

D being a residue of an amine containing biologically active molecule and

A being a leaving group;

X is a spacer moiety such as R5-Y6

Y1, Y2 can each be either O, S, or NR6, independently of each other.

Y3, Y5 can each be either O or S, independently of each other.

Y4 is O, NR6, or -C(R7)(R8)-

Y6 is O, S, NR6, succinimide, maleimide, unsaturated carbon-carbon bonds or any heteroatom containing a free electron pair or is absent.

R2 and R3 are selected independently from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-



substituted heteroaryls, cyano, nitro, halogen, carboxy, carboxyalkyl, alkylcarbonyl, or carboxamidoalkyl;

R4 is selected independently from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl, substituted or non-substituted linear, branched, or cyclical alkoxy, substituted or non-substituted linear, branched, or cyclical heteroalkyloxy, aryloxy, or heteroaryloxy, cyano, halogen;

R5 is selected from substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryls;

R7 and R8 are selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryls, carboxyalkyl, alkylcarbonyl, carboxamidoalkyl, cyano, or halogen;

R6 is selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryls;

R1 is a polymer;

W is selected from substituted or non-substituted linear, branched or cyclical alkyl, aryls, substituted aryls, substituted or non-substituted linear, branched or cyclical heteroalkyl, substituted or nonsubstituted heteroaryls;

Nu is a nucleophile;

n is zero or a positive integer; and

Ar is a multi-substituted aromatic hydrocarbon or a multi-substituted aromatic heterocycle.

4. The prodrug of one of claims 1 to 3, wherein the biologically active moiety is selected from the group of biologically moieties consisting of small molecule biologically active agents or biopolymers.

5. The prodrug of claim 4, wherein the biopolymers are selected from the group of biopolymers consisting of proteins, polypeptides, oligonucleotides and peptide nucleic acids.

6. The prodrug of claim 5, wherein the polypeptides are selected from the group of polypeptides consisting of ACTH, adenosine deaminase, agalsidase, albumin, alfa-1 antitrypsin (AAT), alfa-1 proteinase inhibitor (API), alteplase, anistreplase, anicard serine protease, antibodies (monoclonal or polyclonal, and fragments or fusions), antithrombin III, antitrypsins, aprotinin, asparaginases, biphalin, bone-morphogenic proteins, calcitonin (salmon), collagenase, DNase, endorphins, enfuvirtide, enkephalins, erythropoietins, factor VIIa, factor VIII, factor VIIa, factor IX, fibrinolysin, fusion proteins, follicle-stimulating hormones, granulocyte colony stimulating factor (G-CSF), galactosidase, glucagon, glucagon-like peptides like GLP-1, glucocerebrosidase, granulocyte macrophage colony stimulating factor (GM-CSF), phospholipase-activating protein (PLAP), gonadotropin chorionic (hCG), hemoglobins, hepatitis B vaccines, hirudin, hyaluronidases, iduronidase, immune globulins, influenza

5 vaccines, interleukins (1 alfa, 1 beta, 2, 3, 4, 6, 10, 11, 12), IL-1 receptor antagonist (rhIL-1ra),  
 insulins, interferons (alfa 2a, alfa 2b, alfa 2c, beta 1a, beta 1b, gamma 1a, gamma 1b),  
 keratinocyte growth factor (KGF), transforming growth factors, lactase, leuprolide,  
 levothyroxine, luteinizing hormone, lyme vaccine, natriuretic peptide, pancrelipase, papain,  
 10 parathyroid hormone, PDGF, pepsin, platelet activating factor acetylhydrolase (PAF-AH),  
 prolactin, protein C, octreotide, secretin, sermorelin, superoxide dismutase (SOD),  
 somatropins (growth hormone), somatostatin, streptokinase, sucrase, tetanus toxin fragment,  
 tilactase, thrombins, thymosin, thyroid stimulating hormone, thyrotropin, tumor necrosis  
 factor (TNF), TNF receptor-IgG Fc, tissue plasminogen activator (tPA), TSH, urate oxidase,  
 15 urokinase, vaccines, and plant protein such as lectin and ricin.

7. The prodrug of claim 5, wherein the protein is a protein prepared by recombinant DNA technology.
- 15 8. The prodrug of claim 5, wherein the protein is selected from the group of proteins consisting of antibody fragments, single chain binding proteins, catalytic antibodies and fusion proteins.
9. The prodrug of claim 5, wherein the protein is selected from the group of proteins consisting of antibodies, calcitonin, G-CSF, GM-CSF, erythropoietins, hemoglobins, interleukins,  
 20 insulins, interferons, SOD, somatropin, TNF, TNF-receptor-IgC Fc, glucagon-like peptides like GLP-1,
10. The prodrug of claim 4, wherein the small molecule biologically active agents are selected from the group of agents consisting of central nervous system-active agents, anti-infective,  
 25 anti-neoplastic, antibacterial, anti-fungal, analgesic, contraceptive, anti-inflammatory, steroidal, vasodilating, vasoconstricting, and cardiovascular agents with at least one primary or secondary amino group.
11. The prodrug of claim 4, wherein the small molecule biologically active agents are selected from the group of compounds consisting of daunorubicin, doxorubicin, idarubicin,  
 30 mitoxantron, aminoglutethimide, amantadine, diaphenylsulfon, ethambutol, sulfadiazin, sulfamerazin, sulfamethoxazol, sulfalen, clinafloxacin, moxifloxacin, ciprofloxacin, enoxacin, norfloxacin, neomycin B, spectinomycin, kanamycin A, meropenem, dopamin, dobutamin, lisinopril, serotonin, acivicin and carbutamid.
- 35 12. The prodrug of claim 2 or 3, wherein R4 is selected from the group of small substituents consisting of hydrogen, methyl, ethyl, ethoxy, methoxy, and other linear alkyls, cycloalkyls or branched alkyls and heteroalkyl comprising one to six carbon atoms.

13. The prodrug of claim 2 or 3, wherein R1 is selected from the group of polymers consisting of polyalkyloxy-based polymers like poly(propylene glycol) or poly(ethylene glycol), dex~~tr~~an, chitosan, hyaluronic acid and derivatives, alginate, xylan, mannan, carrageenan, agarose, cellulose, starch, hydroxyethyl starch (HES) and other carbohydrate-based polymers, poly(vinyl alcohols), poly(oxazolines), poly(anhydrides), poly(ortho esters), poly(carbonates), poly(urethanes), poly(acrylic acids), poly(acrylamides) such as poly(hydroxypropylmethacrylamide) (HMPA), poly(acrylates), poly(methacrylates) like poly(hydroxyethylmethacrylate), poly(organophosphazenes), poly(siloxanes), poly(vinylpyrrolidone), poly(cyanoacrylates), poly(esters) such as poly(lactic acid) or poly(glycolic acids), poly(iminocarbonates), poly(amino acids) such as poly(glutamic acid), collagen, gelatin, copolymers, grafted copolymers, cross-linked polymers, and block copolymers from the above listed polymers.
14. The prodrug of claim 2 or 3, wherein R1 is a hydrogel.
15. The prodrug of claim 2 or 3, wherein R1 is a branched or hyperbranched polymer.
16. The prodrug of claim 2 or 3, wherein R1 is a dendrimer or dense star polymer.
17. The prodrug of claim 2 or 3, wherein R1 is a biopolymer.
18. The prodrug of claim 17, wherein R1 is a protein.
19. The prodrug of claim 18, wherein the protein is albumin, an antibody, fibrin, casein or any other plasma protein.
20. The prodrug of any one of claims 2 to 19, wherein R1 further includes one or more biologically active substances.
21. The prodrug of any one of claims 2 to 20, wherein the polymer of R1 has at least one functional group for linkage to X.
22. The prodrug of claim 21, wherein the at least one functional group is selected from the group of functional groups consisting of carboxylic acid and activated derivatives, amino, maleimide, thiol, sulfonic acid and derivatives, carbonate and derivatives, carbamate and derivatives, hydroxyl, aldehyde, ketone, hydrazine, isocyanate, isothiocyanate, phosphoric acid and derivatives, phosphonic acid and derivatives, haloacetyl, alkyl halides, acryloyl, arylating agents like aryl fluorides, hydroxylamine, disulfides like pyridyl disulfide, vinyl sulfone, vinyl ketone, diazoalkanes, diazoacetyl compounds, epoxide, oxirane, and aziridine.

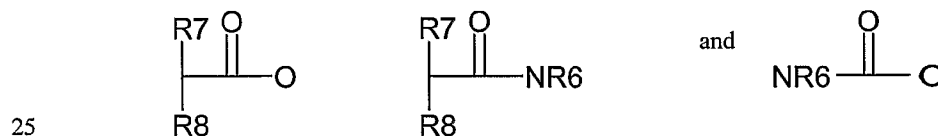
23. The prodrug of claim 21 or 22, wherein the at least one functional group is selected from the group of functional groups consisting of thiol, maleimide, amino, carboxylic acid and derivatives, carbonate and derivatives, carbamate and derivatives, aldehyde, and haloacetyl.

24. The prodrug of one of claims 21 to 23, wherein the bond or group formed between X and R1 is selected from the group of bonds or groups consisting of disulfide, S-succinimido, amide, amino, carboxylic ester, sulphonamide, carbamate, carbonate, ether, oxime, hydrazone, urea, thiourea, phosphate, phosphonate.

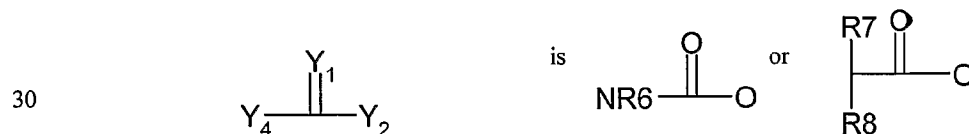
25. The prodrug of any one of claims 21 to 24, wherein the bonds or groups formed between X and R1 is selected from the group of bonds or groups consisting of S-succinimido, amide, carbamate, and urea.

26. The polymeric cascade prodrug linker reagent of any one of claims 2 to 25, wherein A is selected from the group of leaving groups consisting of chloride, bromide, fluoride, nitrophenoxy, imidazolyl, N-hydroxysuccinimidyl, N-hydroxybenzotriazolyl, N-hydroxyazobenzotriazolyl, pentafluorophenoxy and N-hydroxysulfosuccinimidyl.

27. The prodrug of claim 2 or 3, wherein the moiety  $\text{Y}_4 - \text{C}(=\text{Y}_1) - \text{Y}_2$  is preferably selected from the group of moieties consisting of

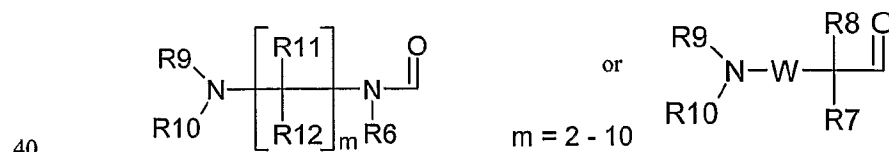


28. The prodrug of claim 27, wherein the moiety



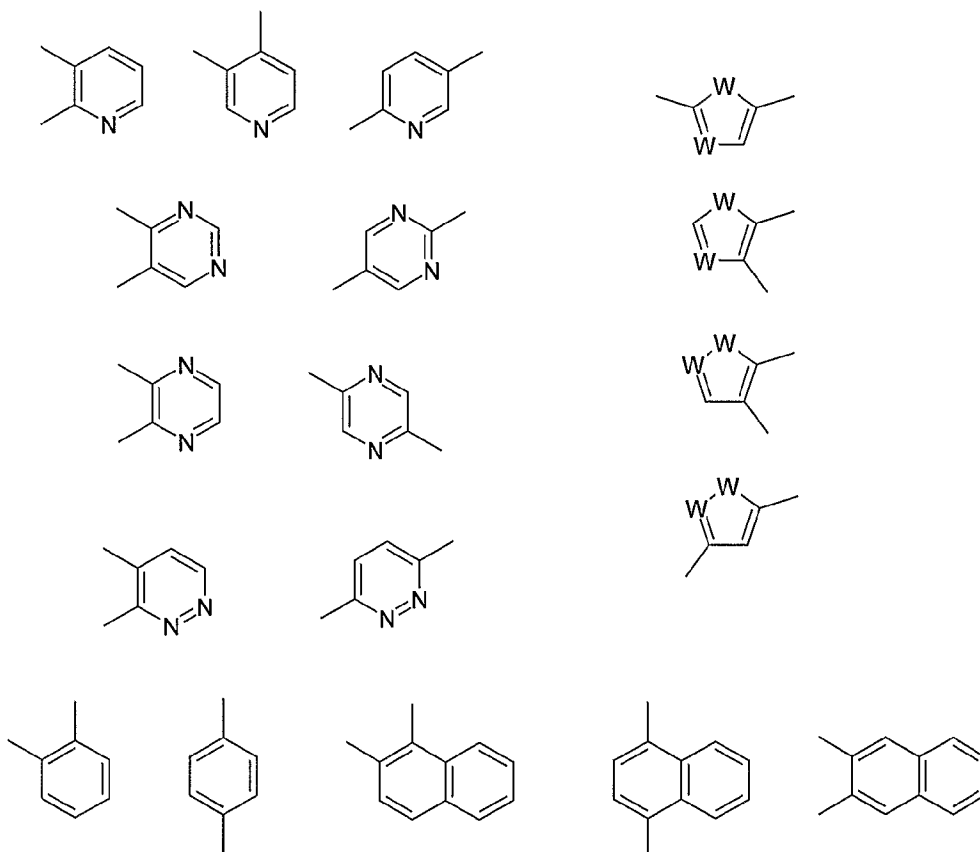
29. The prodrug of claim 27 or 28 wherein R6 may be a further Nu-W.

30. The prodrug of any one of claims 2 to 29 wherein  $\text{Nu-W-Y}_4 - \text{C}(=\text{Y}_1)$  is



in which R9, R10, R11 and R12 are selected independently from hydrogen, substituted or non-substituted alkyl or heteroalkyl, substituted or non-substituted aryl or heteroaryl.

- 5 31. The prodrug of claim 30, wherein R9, 10, R11 and R12 are selected independently from hydrogen or substituted or non-substituted alkyl.
32. The prodrug of any one of claims 2 to 31, wherein Nu is selected from the group of nucleophiles consisting of primary, secondary and tertiary amino groups, thiol, carboxylic  
10 acid, hydroxylamine, hydrazine and nitrogen containing heteroaryl.
33. The prodrug of claim 30 or 31, wherein R7 and/or R8 are not hydrogen.
34. The prodrug of claim 2 or 3, wherein Ar is selected from the group of moieties having the  
15 following structure:



and wherein W is O, S, or N, independent from each other.

35. The prodrug of claim 2 or 3 wherein Ar is a monocyclic or bicyclic aromatic hydrocarbon or aromatic heterocycle.

36. The prodrug of claim 2 or 3, wherein the Ar is a five-membered or six-membered aromatic hydrocarbon or aromatic heterocycle.

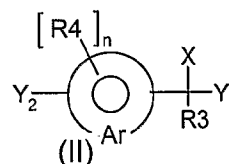
37. The polymeric cascade prodrug linker reagent of any one of the above claims 2-36, in which T is a leaving group A, for covalent conjugation with a biologically active moiety .

38. The polymeric cascade prodrug linker reagent of claim 37, wherein the biologically active moiety is an amine-containing molecule.

39. The polymeric cascade prodrug linker reagent of claim 37 or 38, wherein the biologically active moiety is a peptide, polypeptide or protein.

40. Method for the synthesis of a polymeric prodrug comprising:

- providing a starting molecule of Formula II



- synthesising at least one intermediate compound from the starting molecule of Formula II; and

- attaching an amine-containing biologically active moiety D to the at least one intermediate compound to form the polymeric prodrug;

wherein

Y<sub>2</sub> is selected from O, S, or NR<sub>6</sub>

Y<sub>3</sub> is selected from O or S

X is a spacer moiety such as R<sub>5</sub>-Y<sub>6</sub>;

R<sub>3</sub> is selected independently from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl, cyano, nitro, halogen, carboxy, carboxyalkyl, alkylcarbonyl or carboxamidoalkyl;

R<sub>4</sub> is selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl, substituted or non-substituted linear, branched, or cyclical alkoxy, substituted or non-substituted linear, branched, or cyclical heteroalkyloxy, aryloxy or heteroaryloxy, cyano, or halogen;

R<sub>5</sub> is selected from substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryls ;

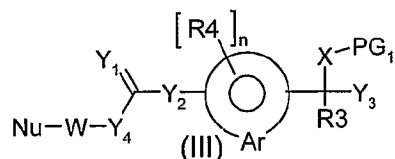
R<sub>6</sub> is selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl; and

Y<sub>6</sub> is O, S, NR<sub>6</sub>, succinimide, maleimide, unsaturated carbon-carbon bonds or a heteratom containing a free electron pair;

n is zero or a positive integer and

Ar is a multi-substituted aromatic hydrocarbon or a multi-substituted aromatic heterocycle.

41. The method of claim 40, wherein a first one of the at least one intermediate compounds is an intermediate molecule of Formula III



synthesised by acylating Y<sub>2</sub> with  $\text{Nu}-\text{W}-\text{Y}_4-\text{C}(=\text{Y}_1)-\text{Y}_2$

and optionally attaching a first protecting group PG<sub>1</sub>,

wherein Nu is a nucleophile;

W is selected from substituted or non-substituted linear, branched or cyclical alkyl, aryls, substituted aryls, substituted or non-substituted linear, branched or cyclical heteroalkyl, substituted or nonsubstituted heteroaryls;

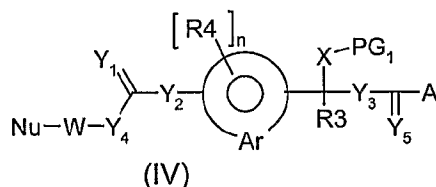
Y<sub>1</sub> is selected from the group of moieties consisting of O, S, or NR<sub>6</sub>;

Y<sub>4</sub> is selected from the group of moieties consisting of -C(R<sub>7</sub>)(R<sub>8</sub>)-, O, or NR<sub>6</sub>;

R<sub>7</sub> and R<sub>8</sub> are selected independently from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl, carboxyalkyl, alkylcarbonyl, carboxamidoalkyl, cyano or halogen.

R<sub>6</sub> is selected selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryls, substituted aryls, substituted or non-substituted heteroaryls;

42. The method of claim 41, wherein a second one of the at least one intermediate compound is an intermediate compound of formula IV



formed by activating the compound of Formula III with an activating agent, wherein A is a leaving group and Y<sub>5</sub> is selected from O or S.

43. The method of claim 42, wherein A is selected from chloride, bromide, fluoride, nitrophenoxy, imidazolyl, N-hydroxysuccinimidyl, N-hydroxybenzotriazolyl, N-hydroxyazobenzotriazolyl, pentafluorophenoxy and N-hydroxysulfosuccinimidyl.

5 44. The method of one of claims 42 or 43, wherein the activating agent is selected from the group of activating agents consisting of 4-nitrophenyl chloroformate or disuccinyl carbonate.

45. The method of one of claims 42 or 43, wherein the amine-containing biologically active moiety D is attached to the second one of the at least one intermediary compound (Formula IV) by displacement of the leaving group A.

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46. The method of claim 45 comprising a step of removal of the reversible first protecting group PG<sub>1</sub> from the second one of the at least one intermediate compound.

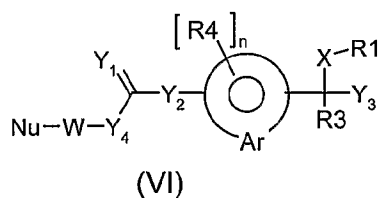
15 47. The method of claim 46, wherein the step of removal of the reversible protection group PG<sub>1</sub> is carried out using a reagent selected from the group of reagents consisting of trifluoroacetic acid or DTT.

48. The method of any one of claims 46 or 47, further comprising a step of attaching a polymer R1 to X.

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49. The method of claim 41, further comprising a step of removing the first polymer protection group PG<sub>1</sub> and attaching a polymer R1 to X to form a third one of the at least one intermediate compound and having a Formula VI.

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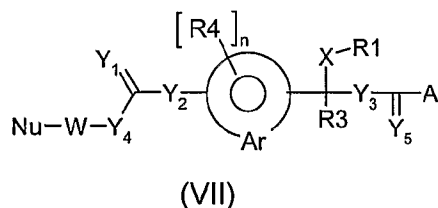
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50. The method of claim 49, further comprising a step of activating the third one of the at least one intermediate compound by using an activating agent to form a fourth one of the at least one intermediate compound and having a Formula VII and wherein A is a leaving group.

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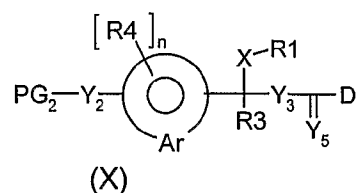


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51. The method of claim 50, wherein A is selected from chloride, bromide, fluoride, nitrophenoxy, imidazolyl, N-hydroxysuccinimidyl, N-hydroxybenzotriazolyl, N-hydroxyazobenzotriazolyl, pentafluorophenoxy and N-hydroxysulfosuccinimidyl.
52. The method of claim 50 or 51, wherein the activating agent is selected from the group of activating agents consisting of 4-nitrophenyl chloroformate or disuccinyl carbonate.
53. The method of one of claim 50 to 52, wherein the amine-containing biologically active moiety D is attached to the compound of Formula VII by displacement of the leaving group A.
54. The method of claim 40, further comprising a step of reacting the starting molecule of Formula II with a first polymer R1 to attach the first polymer R1 to X.
55. The method of claim 54, further comprising a step of protecting Y<sub>2</sub> with a second protective group PG<sub>2</sub>.
56. The method of claim 54 or 55, further comprising the step of activating the prodrug with an activating agent and reacting with an amine-containing biologically active moiety D to form a fifth one of the at least one intermediate compound and having Formula X

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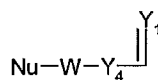
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57. The method of claim 56, wherein the activating agent is selected from the group of activating agents consisting of 4-nitrophenyl chloroformate or disuccinyl carbonate
58. The method of claim 56 or 57, wherein the second protecting group PG<sub>2</sub> is removed from Y<sub>2</sub> and Y<sub>2</sub> is acylated with



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wherein W is selected from substituted or non-substituted linear, branched or cyclical alkyl, aryls, substituted aryls, substituted or non-substituted linear, branched or cyclical heteroalkyl,

substituted or nonsubstituted heteroaryls ;

Nu is a nucleophile;

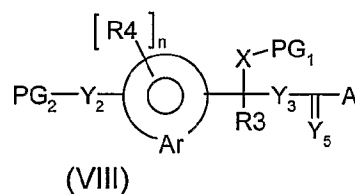
$Y_1$  is selected from the group of moieties consisting of O, S, or  $NR_6$ ;

$Y_4$  is selected from the group of moieties consisting of  $-C(R_7)(R_8)-$ , O,  $NR_6$ ; and

$R_7$  and  $R_8$  are selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl, carboxyalkyl, alkylcarbonyl, carboxamidoalkyl, cyano or halogen.

59. The method of claim 58, wherein the removal of the second reversible group  $PG_2$  is carried out using a reagent selected from the group of reagents consisting of trifluoroacetic acid or DTT.

60. The method of claim 40 comprising a step of attaching a removable first protecting group  $PG_1$  to X and a removable second protecting group  $PG_2$  to the starting compound of Formula II and thereafter activation using an activating agent to form a sixth one of the at least one intermediate compound and having Formula VIII, wherein A is a leaving group



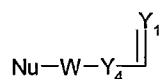
61. The method of claim 60, wherein A is selected from the group of leaving groups selected from the group of leaving groups consisting of chloride, nitrophenoxy, imidazolyly, N-hydroxysuccinimidyl, N-hydroxybenzotriazolyl, N-hydroxyazobenzotriazolyl, pentafluorophenoxy and N-hydroxysulfosuccinimidyl.

62. The method of claim 60 or 61, wherein the activating agent is selected from the group of activating agents consisting of 4-nitrophenyl chloroformate or disuccinyl carbonate.

63. The method of any one of claims 60 to 62, wherein the amine-containing biologically active moiety D is attached to the sixth one of the at least one intermediate compound by displacement of the leaving group A.

64. The method of claim 63 comprising a step of removing the first protecting group  $PG_1$  from X and attaching a polymer R1.

65. The method of claim 64, wherein the second protecting group  $PG_2$  is removed from  $Y_2$  and  $Y_2$  is acylated with



wherein W is selected from substituted or non-substituted linear, branched or cyclical alkyl, aryls, substituted aryls, substituted or non-substituted linear, branched or cyclical heteroalkyl, substituted or nonsubstituted heteroaryls;

Nu is a nucleophile;

5 Y1 is selected from the group of moieties consisting of O, S, or NR6;

Y4 is selected from the group of moieties consisting of -C(R7)(R8)-, O, or NR6;

and

10 R7 and R8 are selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyl or heteroalkyl, aryl, substituted aryl, substituted or non-substituted heteroaryl, carboxyalkyl, alkylcarbonyl, carboxamidoalkyl, cyano or halogen.

66. The method of claim 63, wherein the second protecting group PG<sub>2</sub> is removed from Y<sub>2</sub> and Y<sub>2</sub> is acylated with



wherein W is selected from substituted or non-substituted linear, branched or cyclical alkyl, aryls, substituted aryls, substituted or non-substituted linear, branched or cyclical heteroalkyl, substituted or nonsubstituted heteroaryls;

Nu is a nucleophile;

20 Y1 is selected from the group of moieties consisting of O, S, or NR6;

Y4 is selected from the group of moieties consisting of -C(R7)(R8)-, O, or NR6; ; and

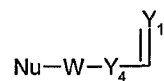
R7 and R8 are selected from hydrogen, substituted or non-substituted linear alkyl, branched alkyl or cyclical alkyl, aryl, substituted aryl, substituted or non-substituted heteroalkyl or heteroaryl, carboxyalkyl, alkylcarbonyl, carboxamidoalkyl, cyano or halogen.

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67. The method of claim 66 comprising a step of removing the first protecting group PG<sub>1</sub> from X and attaching a polymer R1.

68. The method of claim 54 comprising a step acylating Y<sub>2</sub> with

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to form a third one of the at least one intermediate compound;

35 wherein W is selected from substituted or non-substituted linear, branched or cyclical alkyl, aryls, substituted aryls, substituted or non-substituted linear, branched or cyclical heteroalkyl, substituted or nonsubstituted heteroaryls ;

Nu is a nucleophile;

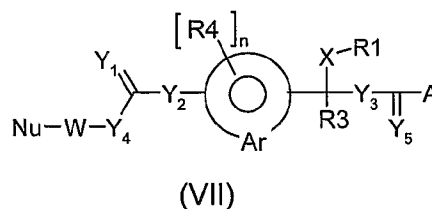
Y<sub>1</sub> is selected from the group of moieties consisting of O, S, or NR6;

40 Y<sub>4</sub> is selected from the group of moieties consisting of -C(R7) (R8)-, O or NR6; and

R7 and R8 are selected from hydrogen, substituted or non-substituted linear alkyl, branched alkyl or cyclical alkyl, aryl, substituted aryl, substituted or non-substituted heteroalkyl or heteroaryl, carboxyalkyl, alkylcarbonyl, carboxamidoalkyl, cyano or halogen.

- 5 69. The method of claim 68, further comprising a step of activating the third one of the at least one intermediate compound by using an activating agent to form a seventh one of the at least one intermediate compound and having a Formula VII and wherein A is a leaving group

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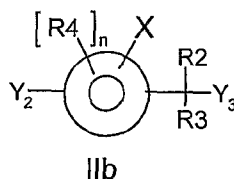
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70. The method of claim 69, wherein A is selected from the group of leaving groups consisting of chloride, nitrophenoxy, imidazolyl, N-hydroxysuccinimidyl, N-hydroxybenzotriazolyl, N-hydroxyazobenzotriazolyl, pentafluorophenoxy and N-hydroxysulfosuccinimidyl.

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71. The method of claim 69 or 70, wherein the activating agent is selected from the group of activating agents consisting of 4-nitrophenyl chloroformate or disuccinyl carbonate.
72. The method of one of claims 69 to 71, wherein the amine-containing biologically active moiety D is attached to the compound of Formula VII by displacement of the leaving group A.

73. The method of any one of claims 40 to 72 wherein the starting molecule of Formula II is replaced by a starting molecule of Formula IIb



wherein R2 is selected from hydrogen, substituted or non-substituted linear, branched or cyclical alkyls or heteroalkyls, aryls, substituted or non-substituted heteroaryl, cyano, nitro, halogen, carboxy, carboxyalkyl, alkylcarbonyl or carboxamidoalkyl.

74. A method for hydrolysing the prodrug of any one of claims 1 to 36 comprising a step of placing the prodrug in solution with a pH of approximately 7.4
75. The method of claim 74, wherein the solution is an extra-cellular fluid.
76. Method of administration of an amine-containing moiety to a living organism comprising:
- a first step of providing a polymeric cascade prodrug according to any one of claims 1 to 39;
  - a second step of administering the polymeric cascade prodrug to the living organism; and
  - a third step of cleaving the amine-containing moiety from the polymeric cascade prodrug by means of a substantially non-enzymatic reaction.
77. The method of claim 76, wherein the third step is carried out in an extra-cellular fluid.
78. The method of one of claims 76 or 77, wherein the substantially non-enzymatic reaction comprises a step of hydrolysis.
79. The method of one of claims 76 to 78, wherein the substantially non-enzymatic reaction comprises a step of intramolecular cyclization or intramolecular catalysis.
80. The method of one of claims 76 to 79, further comprising a step of sterically protecting at least part of the linker by a sterically demanding carrier.

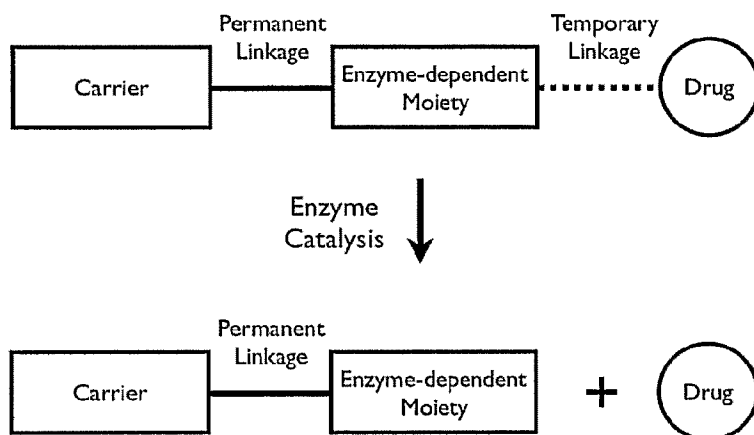
81. In an polymeric cascade prodrug according to any one of claims 1 to 39, a method of cleaving the amine-containing moiety from the carrier by a substantially non-enzymatic reaction of the nucleophile-containing linker.
- 5 82. The method of claim 81, wherein the substantially non-enzymatic reaction is carried out at a pH of approximately 7.4.
83. The method of claim 81 or 82, wherein the amine-containing moiety attached to the carrier is cleaved in an extra-cellular fluid.
- 10 84. The method of one of claims 81 to 83, wherein the substantially non-enzymatic reaction comprises a step of hydrolysis.
85. The method of one of claims 81 to 84, wherein the substantially non-enzymatic reaction comprises a step of intramolecular cyclization or catalysis.
- 15 86. The method of one of claims 81 to 85, further comprising a step of sterically protecting at least part of the nucleophile-containing linker by a sterically demanding carrier.
- 20 87. The method of any one of claims 81 to 86, wherein the amine-containing moiety is a biologically active moiety.
88. A method of providing a therapeutically useful concentration of a biologically active molecule by *in vivo* cleavage of the biologically active molecule from the prodrug according to any one of claims 1 to 39.
- 25

89. Fig. 1



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Fig. 2



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Fig. 3

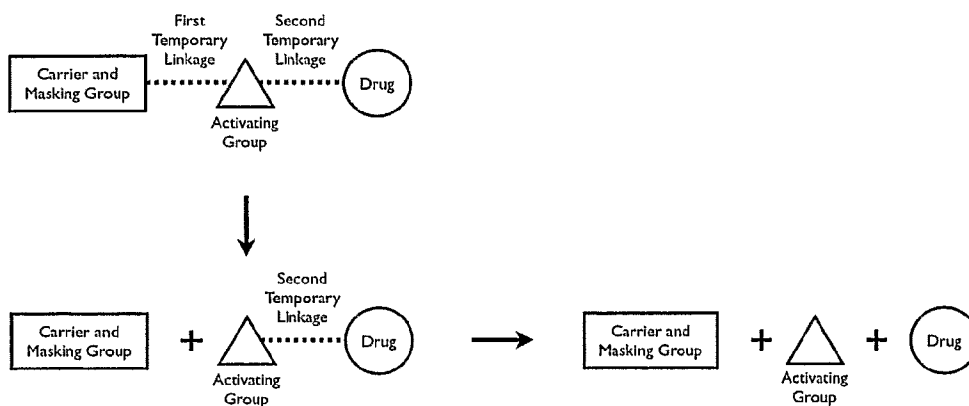


Fig. 4

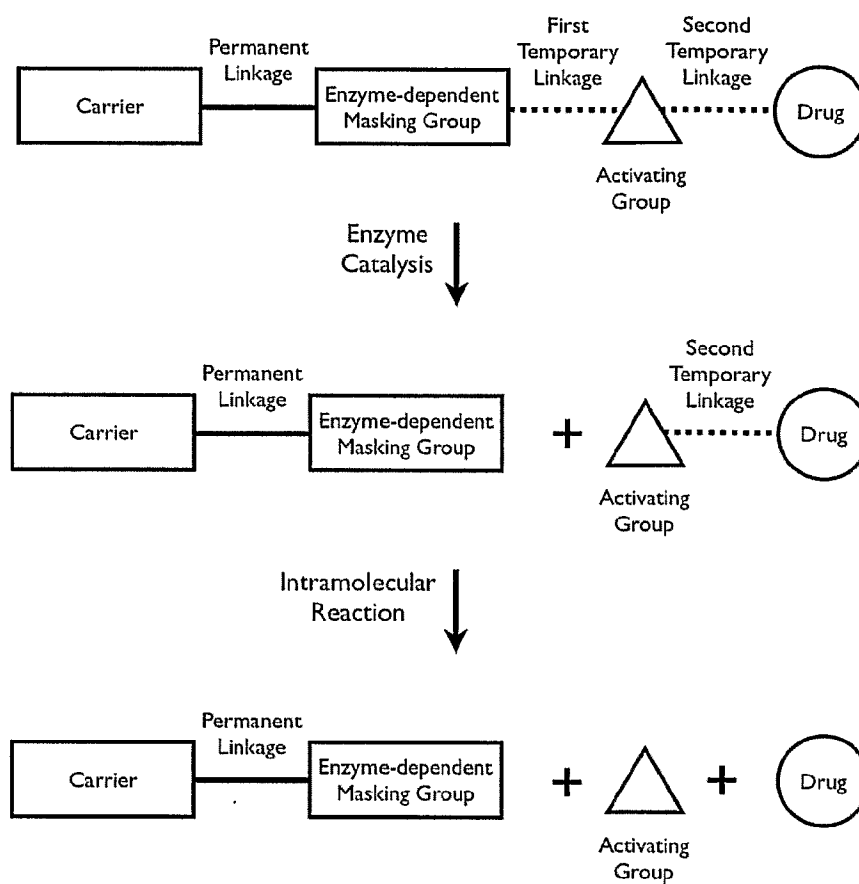




Fig. 5

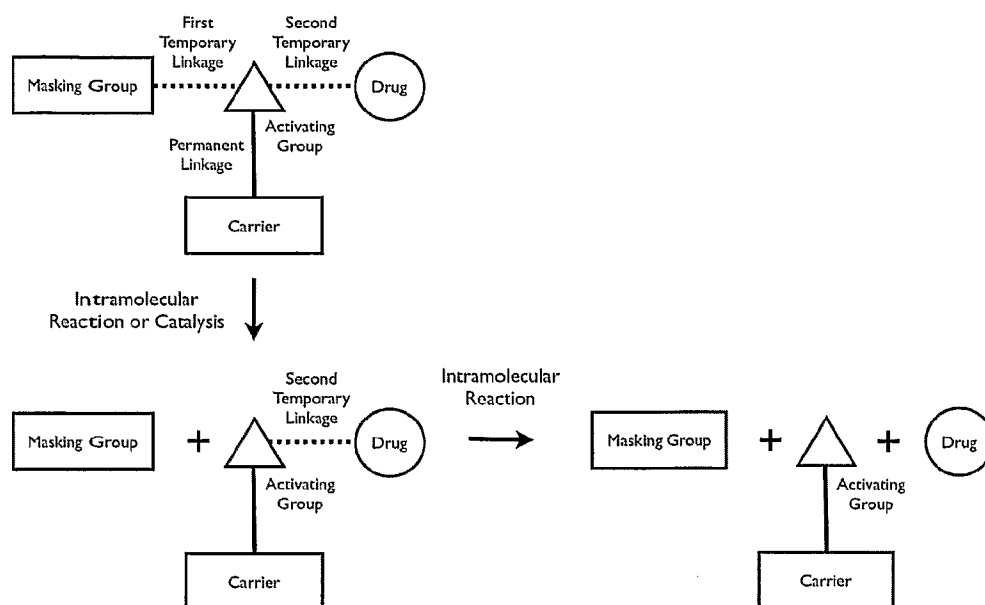
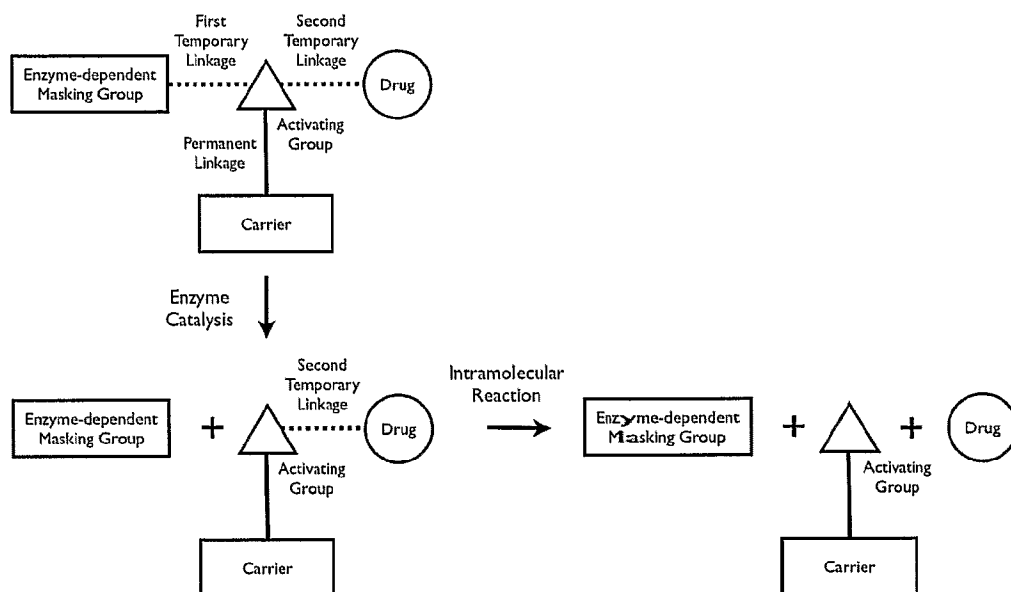
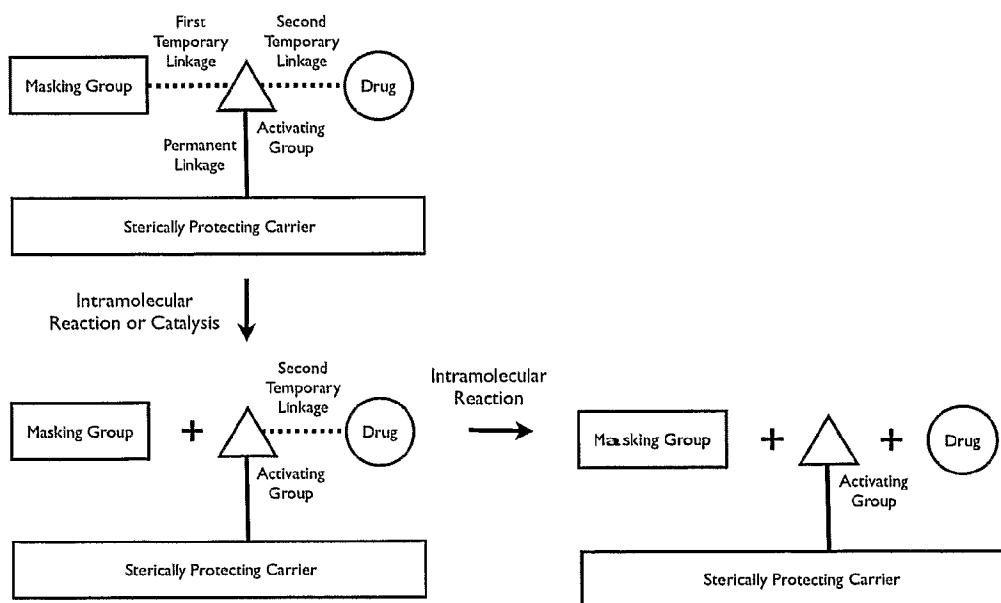


Fig. 6



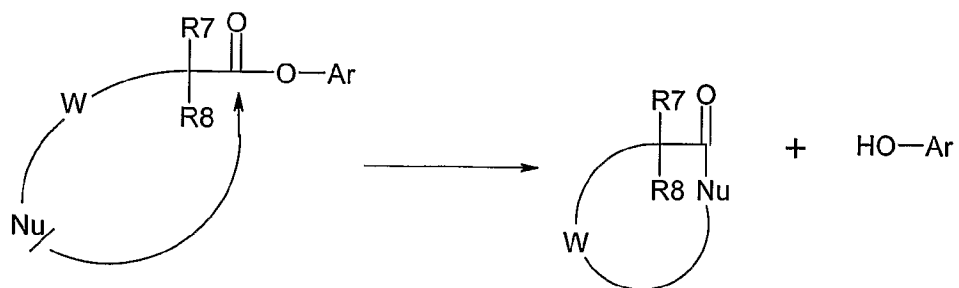
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Fig. 7



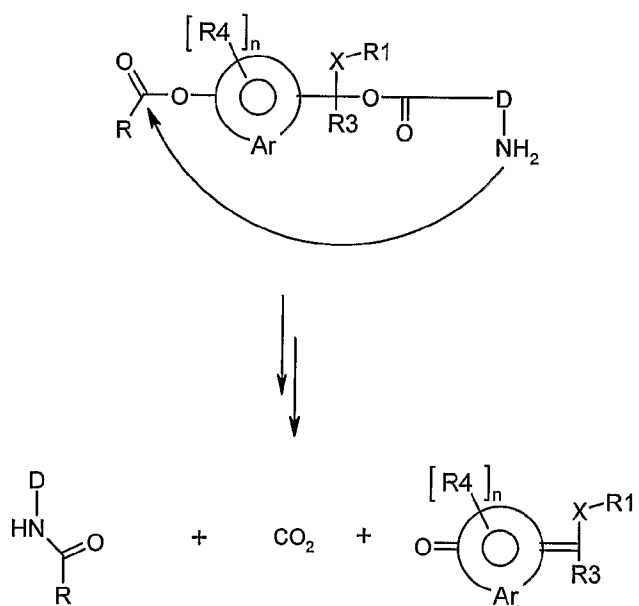
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Fig. 8



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Fig. 9



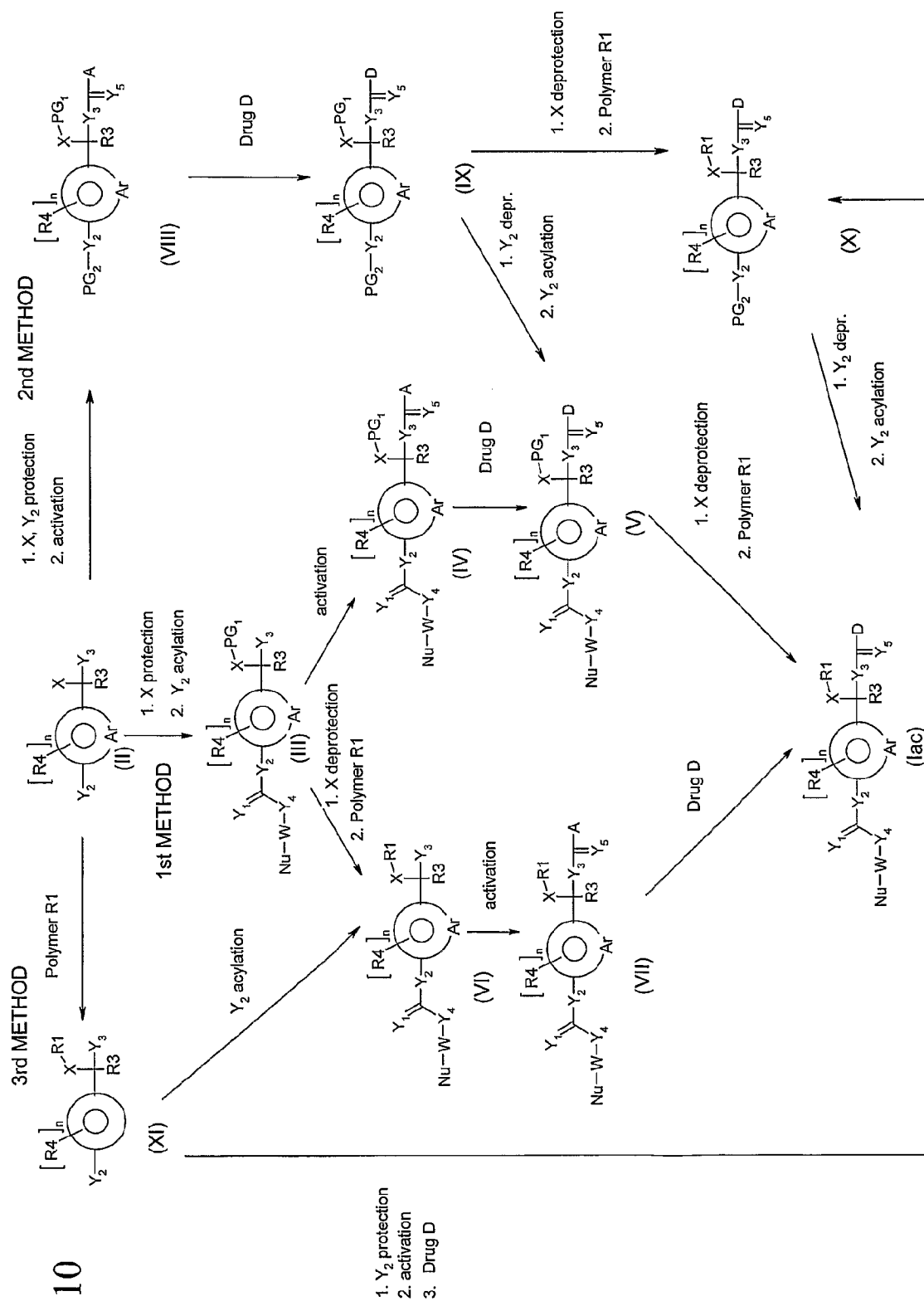
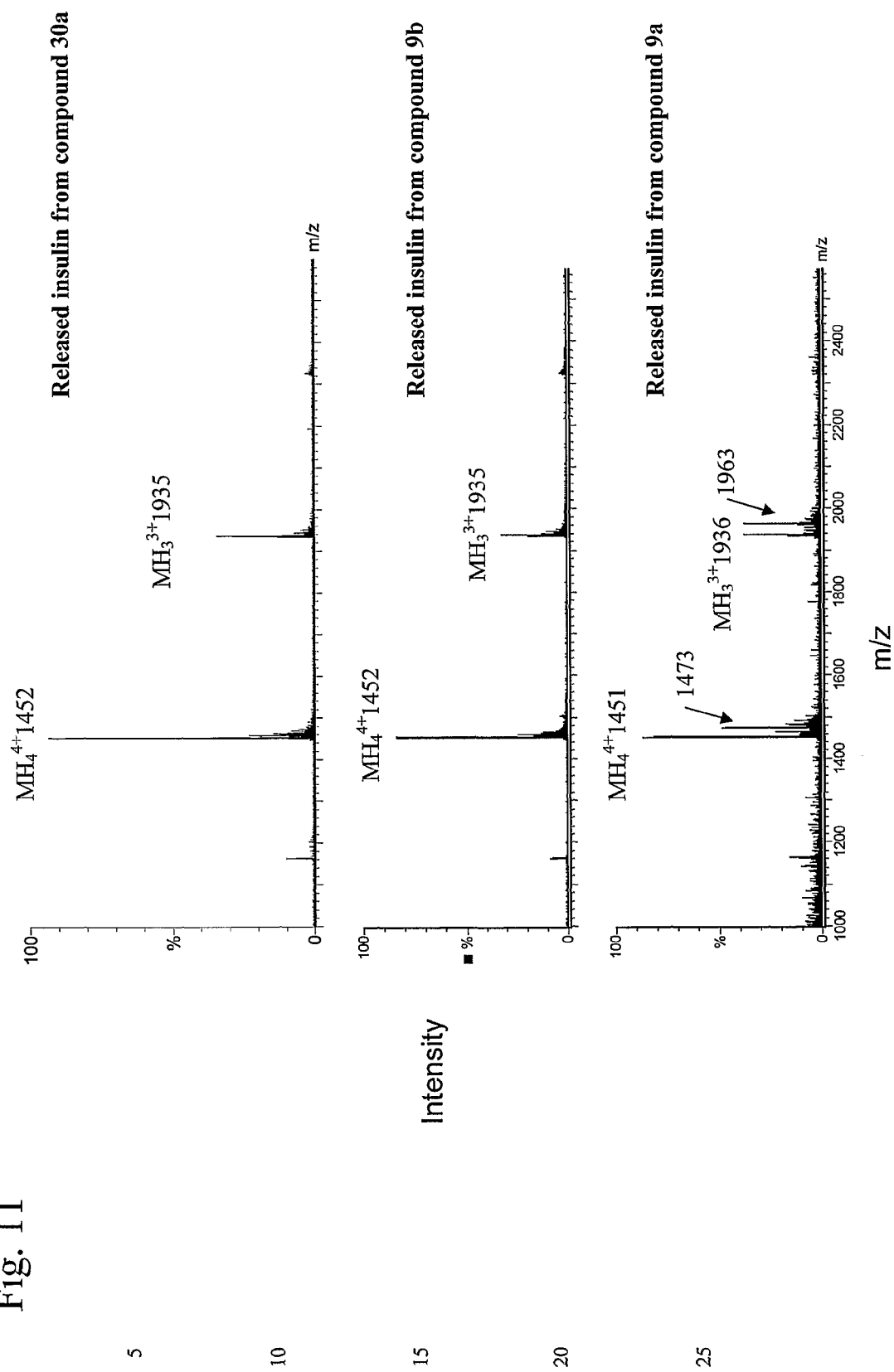


Fig. 11



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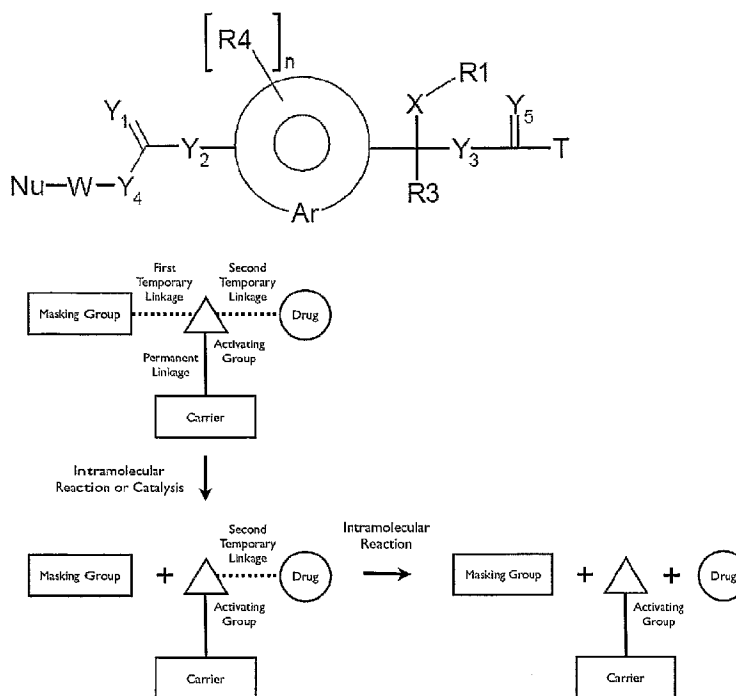
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(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI,

[Continued on next page]

(54) Title: POLYMERIC PRODRUG WITH A SELF-IMMOBILIZATIVE LINKER



(57) Abstract: A cascade carrier linked prodrug is described which comprises a biologically active moiety and - a masking group having at least one nucleophile and being distinct from the carrier. In one preferred embodiment of the invention, the prodrug has the structure represented by formula (I).



FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO,  
SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN,  
GQ, GW, ML, MR, NE, SN, TD, TG).

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**Published:**

— with international search report

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/EP2005/003061

A. CLASSIFICATION OF SUBJECT MATTER  
INV. A61K47/48

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, BIOSIS, CHEM ABS Data, EMBASE

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 03/057716 A (BOERTH NANCY JOHNSTON ; NEW RIVER PHARMACEUTICALS INC (US); BISHOP BAR) 17 July 2003 (2003-07-17)  page 7 - page 8; claims 10,11; figures 3-5 page 5, paragraph 2	1,2,4, 10-13, 21-25, 32, 34-36, 74-88
X	WO 02/083180 A (BEUSKER PATRICK HENRY ; BUSSCHER GUUSKE FREDERIKE (NL); SCHEEREN JOHAN) 24 October 2002 (2002-10-24) cited in the application  page 10, lines 25-28; claims 13,14; figures 3,12	1,2,4, 10-12, 20-25, 32-36, 40-72, 74-88

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

7 February 2006

Date of mailing of the international search report

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## INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP2005/003061

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>WO 2004/019993 A (LIST BENJAMIN ; PESSAH NETA (IL); SHAMIS MARINA (IL); AMIR ROEY JACOB) 11 March 2004 (2004-03-11) cited in the application</p> <p>page 5; figure 3; compound IB page 25, line 12 - page 26, line 10 page 32, lines 6-16; claims 3,7,13,16,24,123,129,161</p> <p>-----</p>	<p>1,2,4, 10-12, 20-25, 27-36, 40-72, 74-88</p>
X	<p>SHABAT D. ET AL: "Chemical adaptor systems" CHEM. EUR. J., vol. 10, 22 March 2004 (2004-03-22), XP002298470 cited in the application see conclusions abstract; figures 1,2,8</p> <p>-----</p>	<p>1,2,4, 10-12, 20-25, 27-36, 40-72, 74-88</p>
P,X	<p>WO 2004/043493 A (BEUSKER PATRICK HENRY ; SCHEEREN JOHANNES WILHELM (NL); SYNTARGA B V ( ) 27 May 2004 (2004-05-27) cited in the application</p> <p>page 34, lines 18-20; claims 42,49 page 41, line 10 - page 42, line 28 page 50, line 29 - page 51, line 25 page 12, paragraph 3 page 14, paragraphs 1,2</p> <p>-----</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>
Y	<p>AMIR R. J. ET AL: "Self immolative dendrimers" ANGEW. CHEM. INT., vol. 42, 2003, pages 4494-4499, XP008035926</p> <p>abstract; figure 1</p> <p>-----</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>
Y	<p>SAUERBREI B. ET AL: "An enzyme labile linker group for organic syntheses on solid supports" ANGEW. CHEM. INT. ED., vol. 37, 1998, pages 1143-1146, XP001120792 page 1144, paragraph 2; figure 1</p> <p>-----</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>
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## INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP2005/003061

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>GROOT DE FRANCISCUS M H ET AL: "Elongated Multiple Electronic Cascade and Cyclization Spacer Systems in Activatable Anticancer Prodrugs for Enhanced Drug Release"</p> <p>JOURNAL OF ORGANIC CHEMISTRY, AMERICAN CHEMICAL SOCIETY. EASTON, US, vol. 66, 2001, pages 8815-8830, XP002212035</p> <p>ISSN: 0022-3263</p> <p>abstract; figure 1</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>
A	<p>GREENWALD R B ET AL: "Drug delivery systems employing 1,4- or 1,6-elimination: poly(ethylene glycol) prodrugs of amine-containing compounds"</p> <p>JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY. WASHINGTON, US, vol. 42, September 1999 (1999-09), pages 3657-3667, XP002184836</p> <p>ISSN: 0022-2623</p> <p>cited in the application</p> <p>figures 2,8</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>
A	<p>ANTCZAK C ET AL: "A NEW ACIVICIN PRODRUG DESIGNED FOR TUMOR-TARGETED DELIVERY"</p> <p>BIOORGANIC &amp; MEDICINAL CHEMISTRY, ELSEVIER SCIENCE LTD, GB, vol. 9, 2001, pages 2843-2848, XP001150714</p> <p>ISSN: 0968-0896</p> <p>cited in the application</p> <p>abstract</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>
X	<p>WO 03/026577 A (SEATTLE GENETICS, INC; SENTER, PETER, D; TOKI, BRIAN, E)</p> <p>3 April 2003 (2003-04-03)</p> <p>page 16, lines 8-31</p> <p>pages 6,7; claims 37-44,79,93</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>
E	<p>EP 1 525 890 A (COMPLEX BIOSYSTEMS GMBH)</p> <p>27 April 2005 (2005-04-27)</p> <p>page 69</p> <p>page 71 - page 72</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>

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# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/EP2005/003061

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E	<p>WO 2005/082023 A (GENENTECH, INC; SEATTLE GENETICS, INC; FENG, BAINIAN) 9 September 2005 (2005-09-09)</p> <p>abstract; claims 4,5,34,35; figure 1 -----</p>	<p>1,2,4, 10-13, 20-25, 27-36, 40-72, 74-88</p>

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/EP2005/003061

## Box II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:  
  
Although claims 76-88 are directed to a method of treatment of the human/animal body (Article 52(4) EPC), the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

see annex

### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

## FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. claims: 1,2, 4, 10-13, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a polymer as recited in claim 13 and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 2-31

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2. claims: 1,2, 4-9, 12, 13, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a polymer as recited in claim 13 and T is protein or polypeptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1, 3-31

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3. claims: 1, 2, 4, 5, 12, 13, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a polymer as recited in claim 13 and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1, 2, 4-31

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4. claims: 1, 2, 4, 10-12, 14, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is an hydrogel and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-3, 5-31

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5. claims: 1, 2, 4-9, 12, 14, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a hydrogel and T is a protein or peptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-4, 6-31

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6. claims: 1, 2, 4, 5, 12, 14, 20-25, 27-36, 40-72, 74-88 in part

## FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is an hydrogel and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-5, 7-31

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7. claims: 1, 2, 4, 10-12, 15, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a branched or hyperbranched polymer and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-6, 8-31

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8. claims: 1, 2, 4-9, 12, 15, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a branched or hyperbranched polymer and T is a protein or a peptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-7, 9-31

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9. claims: 1,2, 4, 5, 12, 15, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a branched or hyperbranched polymer and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-8, 10-31

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10. claims: 1, 2, 4, 10-12, 16, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a dendrimer or dense star polymer and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-9, 11-31.

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11. claims: 1, 2, 4-9, 12, 16, 20-25, 27-36, 40-72, 74-88 in part

## FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a dendrimer or dense star polymer and T is a protein or a peptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-10, 12-31

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12. claims: 1, 2, 4, 5, 12, 16, 20-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a dendrimer or dense star polymer and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-11, 13-31

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13. claims: 1, 2, 4, 10-12, 17-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a biopolymer or a protein and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-12, 14-31.

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14. claims: 1,2, 4-9, 12, 17-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a biopolymer or protein and T is a protein or a peptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-13, 15-31

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15. claims: 1, 2, 4, 5, 12, 17-25, 27-36, 40-72, 74-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 2 wherein R1 is a biopolymer or a protein and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-14, 16-31

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16. claims: 1, 3, 4, 10-13, 20-25, 27-36, 40-88 in part

## FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a polymer as recited in claim 13 and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-15, 17-31

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17. claims: 1, 3-9, 12, 13, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a polymer as recited in claim 13 and T is protein or polypeptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-16, 18-31

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18. claims: 1, 3-5, 12, 13, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a polymer as recited in claim 13 and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-17, 19-31

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19. claims: 1, 3, 4, 10-12, 14, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is an hydrogel and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-18, 20-31

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20. claims: 1, 3-9, 12, 14, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a hydrogel and T is a protein or peptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-19, 21-31

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21. claims: 1, 3-5, 12, 14, 20-25, 27-36, 40-88 in part



## FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is an hydrogel and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-20, 22-31

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22. claims: 1, 3, 4, 10-12, 15, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a branched or hyperbranched polymer and T is a biologically active small molecule. Excluding the subject matter of inventions 1-21, 23-31

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23. claims: 1, 3-9, 12, 15, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a branched or hyperbranched polymer and T is a protein or a peptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-22, 24-31

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24. claims: 1, 3-5, 12, 15, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a branched or hyperbranched polymer and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-23, 25-31

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25. claims: 1, 3, 4, 10-12, 16, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a dendrimer or dense star polymer and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-24, 26-31

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26. claims: 1, 3-9, 12, 16, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a dendrimer or dense star polymer and T is a protein or a peptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-25, 27-31

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## FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

27. claims: 1, 3-5, 12, 16, 20-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a dendrimer or dense star polymer and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-26, 28-31

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28. claims: 1, 3, 4, 10-12, 17-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a biopolymer or a protein and T is a biologically active small molecule. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-27, 29-31

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29. claims: 1, 3-9, 12, 17-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a biopolymer or protein and T is a protein or a peptide. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-28, 30-31

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30. claims: 1,3-5, 12, 17-25, 27-36, 40-88 in part

Polymeric cascade prodrug having the structure as depicted by claim 3 wherein R1 is a biopolymer or a protein and T is an oligonucleotide or peptide nucleic acid. Method of administration of the same comprising cleavage from the carrier by means of a non-enzymatic reaction. Excluding the subject matter of inventions 1-29, 31

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31. claims: 26, 37-39 complete; 1-3, 12-25, 27-36, 40-52, 54, 55, 60-62, 64-71 in part

Molecule having the structure as depicted by claims 2 or 3 wherein R1 is a polymer and T is leaving group A. Excluding the subject matter of inventions 1-30

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# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP2005/003061

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